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ASSESSMENT OF ENVIRONMENTAL CONTAMINATION EXPLORATORY STAGE TOOELE ARMY DEPOT TOOELE, UTAH

VOLUME III APPENDICES B THROUGH E

IRP 81-04



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ASSESSMENT OF ENVIRONMENTAL CONTAMINATION VOL

3 OF 5

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ASSESSMENT OF ENVIRONMENTAL CONTAMINATION EXPLORATORY STAGE TOOELE ARMY DEPOT TOOELE, UTAH

VOLUME III APPENDICES B THROUGH E

APPENDIX B: METHODOLOGY FOR PHASE II TECHNICAL EFFORT APPENDIX C: WELL CONSTRUCTION AND LITHOLOGY SUMMARIES

APPENDIX D: GEOPHYSICS PROGRAM

APPENDIX E: CONTAMINATION RESULTS (SITE SUMMARY FORMS)

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MATERIALS AGENCY
ABERDEEN PROVING GROUND, MARYLAND 21010

Prepared by:

KEVIN T. BLOSE ERIC G. LAPPALA ROBERT L. STOLLAR KAREN F. KNIRSCH

ERTEC WESTERN, INC. 3777 LONG BEACH BOULEVARD LONG BEACH, CALIFORNIA 90807



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Appendix B
Methodology for Phase II Technical Effort

B-1 Methodology for Well Installation

B-1.1 Drilling Operations

Monitoring wells for the TEAD Environmental Survey were drilled using hollow stem auger, mud rotary, and cable tool techniques. Drilling contractors were F.M. Fox and Associates, Denver, Colorado for the hollow stem auger and Scott Stephenson Drilling, Fillmore, Utah for the mud rotary and cable tool.

Ten wells and two borings in the South Area were completed using the hollow stem auger; one boring (S-9) was discontinued at a depth of 111 feet, and was capped. One well (N-8A) and one boring (N-2A) in the North Area were drilled with the hollow stem auger.

Drilling procedures were followed as outlined in the Technical Plan. The major exception was that a 6 7/8-inch hole, rather than a 10-inch hole was drilled at most sites. It was determined that if caving were not a problem, it was more efficient to drill a smaller hole, remove the auger, and then install the casing, because the drilling efficiency of the larger auger decreased rapidly at depths greater than 40 feet. If caving was a problem, the hole was drilled with the smaller auger, reamed with the larger auger, and the casing was installed through the hollow stem.

Difficulties forcing bore abandonment were encounterd at three locations: N-2A, S-15, and S-9. Large quartzite boulders and cobbles halted drilling at N-2A and S-15. S-9 was abandoned due to sticky clay and the limit of the auger capabilities. Upon completion of the field program assessment, it was determined that it was unnecessary to complete these wells.

Split spoon samples were taken every 5 feet and were described by the Ertec hydrogeologist in accordance with USATHAMA requirements. Exceptions to the

sampling intervals occurred when drilling through gravels in which there was auger refusal.

Four wells in the North Area were drilled using the mud rotary technique. These wells ranged in depth from 65 feet (N-3B) to 709 feet (N-6). Drilling procedures were followed as outlined in the Technical Plan. Rotary wash samples were obtained over 1-foot intervals every 5 feet, i.e., the hole was circulated with mud for approximately ten minutes after drilling 5 feet, to ensure that most of the cuttings were removed and then the sample was collected during the drilling of the next foot. At 15-foot intervals (the end of each drill rod), the drill stem was removed and a split spoon sample was attempted. Exceptions occurred when the drilling proceeded through gravels, in which case it was impossible to recover a sample. The split spoon sampler was driven using 600 pound cable tool jars. All samples were collected and logged by the Ertec hydrogeologist in accordance with USATHAMA requirements, as set forth in the Ertec field manual and in Appendix C (revised) of the original proposal for this project.

Major drilling problems encountered were lost circulation, caving, and zones consisting of large quartzite boulders and cobbles. Generally, circulation was regained by thickening the mud, but in one case (N-3A), circulation loss was so extreme (thousands of gallons) that it was necessary to use mica flakes as lost circulation material. At N-6, circulation loss and caving were severe enough that rotary drilling could not continue past 30 feet. This necessitated drilling with a cable tool rig and setting surface casing to a depth of 100 feet before drilling was resumed with the rotary rig. At N-7 3600 gallons of mud were lost in the first 4 feet of drilling. N-7 was then also started with the cable tool rig.

Large quartzite boulders and cobbles presented the most challenging problem for the rotary rig. This lithology was previously undocumented at TEAD, except for some shallow holes drilled by the U.S. Army Environmental Hygiene Agency near the North Area Industrial Waste Pond in August, 1981. The depths, extents, and drilling difficulties in these materials were unknown, and therefore unanticipated. These boulders were extremely hard, and there was little fine material between them. Therefore, a natural sieve was created, drilling mud was lost, and drill bits were very quickly worn down. The use of the rotary rig became very inefficient, and it was for this reason that the cable tool rig was employed.

The cable tool rig was used to drill two wells in the North Area (N-2C and N-4). Two wells were started with the cable tool (N-6 and N-7). N-6 was later completed with the rotary rig, and drilling on N-7 was stopped at 75 feet, when this well was deleted from the drilling program. N-2B was drilled to 65 feet where bedrock was penetrated. S-1 was also drilled with the cable tool because flooding at this drill site prohibited the use of the auger rig before the auger program was completed. When access to the drill site was available, the cable tool was the only rig still at TEAD.

Samples of drill cuttings from the bailer used in drilling operations were collected continuously when drilling with the cable tool. Split spoon samples were obtained when feasible by driving the sample with the cable tool jars. Surface casing driven by the cable tool was removed from wells N-2C and N-4. Surface casing was left in N-2B, N-6, and N-7. Numerous attempts to remove the surface casing from N-2B failed. The casing was left in N-6 and N-7 to prevent further caving; it is above the water table in both bores.

B-1.2 Well Construction

Well construction procedures were followed as outlined in the Technical Plan. The major exceptions were the use of 4.5 inch O.D. Schedule 40 polyvinyl chloride (PVC) rather than 4-inch O.D. Schedule 80 PVC and that Well N2C was completed with 2-inch O.D. Schedule 40 PVC casing. Table 1 lists the construction materials, and source or supplier of materials.

N-2C was the only well to be completed as a 2-inch diameter well. It originally had been constructed as a 4.5 inch well, but when well development commenced, the 3.75-inch bailer was unable to proceed past a depth of 20 feet. Upon inspection by the Ertec field hydrogeologist, it was observed that there was a bulge in the PVC, prohibiting the passage of the bailer. This bulge occurred at a flush-threaded joint. It is not known if this was caused by a defect in the PVC. A crack was also observed, therefore a decision was made to recase the well with 2 inch PVC and emplace a second gravel pack and bentonite seal. It was determined that this method was more cost-effective than drilling a new well while still providing an uncompromised sample for chemical analysis.

All lithologic summaries, well construction summaries, and geophysical logs for wells and bores are found in Appendix C.

B-1.3 Well Development

Well development was generally completed following the procedures outlined in the Technical Plan. Wells in the South Area were not drilled with mud, and the aquifer material consisted of very fine grained silts and sands.

Therefore, most of these wells were very particulate-laden, and in many cases the transmissivity was so low that the well could be bailed dry very easily.

Table 1. Well Construction Materials

Material	Brand	Source/Supplier
PVC Casing	4.5 inch O.DSchedule 40, flush-threaded/2.0 inch O.DSchedule 40, flush threaded	Timco Mfg., Prairie du Sac, Wisconsin
PVC Screen	.050 slot, 4.5 inch O.D Schedule 40, flush threaded/ 2.0 inch O.DSchedule 40, flush threaded	Timco Mfg., Prairie du Sac, Wisconsin
Bentonite (drilling fluid)	Aquagel	NL Baroid, Evanston, Wyoming
Bentonite (granular)	Big Horn	Nova Mud, Orem, Utah
Bentonite (pellets)		Nova Mud, Orem, Utah
Gravel/sand pack	8-12 silica sand	Colorado Silica Sand, Colorado Springs, Colorado/ supplier- Vinton Pipe Co., Salt Lake City, Utah
Cement	Portland Type II "Utah Cement"	Ideal Basic Industries, Salt Lake City, Utah

For these wells, the development method consisted of bailing the well dry, allowing full recovery and bailing dry again. This method was repeated a minimum of five times for each well. If it appeared that a well could recover fairly rapidly (a minimum of 2 gallons per minute), a submersible pump was used for further development. The well was then pumped for a minimum of two hours, and pH, Ec, temperature, and sand content were monitored as outlined in the Technical Plan. Table 2 shows the method of development and the amount of water removed from each well.

All wells drilled by the mud rotary techniques were bailed until the well was free of drilling mud. Pump tests were conducted with the use of a submersible pump and appropriate field parameters were recorded.

At Well N-6, the well was continually bailed dry, and it was believed that the mud cake used to prevent lost circulation was too thick to allow for proper well development. Attempts to clean the well using drilling water from existing Well #1 failed, and upon USATHAMA approval, 100 pounds of sodium acid pyrophosphate (SAPP), a dispersing agent, were added to the well. The well was surged repeatedly with a bailer, but most recent water level measurements (June 1982) indicate that the well is dry.

B-2 Methodology for the Collection of Soil, Sediment, and Surface-Water Samples

B-2.1 Soil Samples

Soil samples used for chemical analysis were collected from borings using a split spoon sampler. The spoons were cleaned carefully with a wire brush, rinsed repeatedly in distilled water, and wiped dry with a clean dry rag prior to sampling. Samples were taken in borings installed with the auger by attaching the spoon to an A-rod, lowering it down the center of the hollow

Table 2. Well Development

Well	Development Method	Total Amount of Water Removed (gallons)	Date Development Completed
S-1	pump, 18 gpm	2840	6/1/82
S-2	bail/pump; 1 gpm	482	4/20/82
S-3	bail/pump; 12 gpm	1110	4/18/82
S-4	bail -	216	4/20/82
S-5	bail/pump; 11 gpm	1980	4/18/82
S-6	bail -	104	4/19/82
S-7	bail -	151	4/20/82
S-8	bail -	103	4/20/82
S-10	bail/pump; 4 gpm	842	4/19/82
S-12	bail -	36	4/20/82
S-14	bail/pump; 9 gpm	985	4/18/82
N-2C	pump; 1 gpm	45	6/12/82
N-3A	pump; 15 gpm	3843	6/17/82
N-3B	bail -	46	6/4/82
N-4	<pre>bail/pump; 6 gpm</pre>	770	6/3/82
N-6	bail –	dry	4/20/82
N-8B	bail/pump; 18 gpm	7298	3/2/82

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stem auger and driving it into the soil. Upon removal of the spoon, we carefully inspected the contents, removed the upper 6 inches (or more) of slough, scraped the sides of the sample with a stainless steel blade to remove smear, and placed portions of the sample in a wide-mouth amber glass jar with a Teflon-lined plastic lid. Samples were immediately placed on ice and sent to the lab within 48 hours of collection. Samples were obtained in this manner from Wells S-1, S-2, and S-8 and from Boring S-11. The 11 split spoon samples from Well N-3A were taken with a slightly modified procedure using a down-hole hammer to drive the sample. These borings were drilled by the mud rotary method. Also, greater care was taken in removing the sample from the spoon as drilling mud sometimes contaminated a portion of the sample. The initial sample from Well N-3A was taken as a grab sample. A soil sample from the proposed site of N-1 also was taken as a grab sample. Although N-1 was deleted from the drilling program, the sample was analyzed, and labelled N-1, GS1. Table 1 lists the soil samples collected for chemical analysis.

B-2.2 Sediment Samples

Eight samples of sediment were collected for chemical analysis, four from the TEAD North Area, and four from the TEAD South Area. In the Technical Plan, it was proposed that sediment samples be taken at points immediately downslope from the locations of surface water samples. With the exception of sample N-SD3, which was collected immediately below the location of N-SW2 (the industrial waste pond), it was determined by field personnel that this method was not feasible for the remaining sediment samples as there was no surface water flow at the sample locations.

For the remaining seven locations, the proposed sampling site was inspected by field personnel for fine-grained materials which were likely to have been

Table 1. Soil Samples for Chemical Analysis

Site I.D.	Site Type	Field Sample No.	Sample Depth (Fee	Sample t) Method	Date Collected
N-3A	Well	GS-1	1.0	Grab	2/27/82
N-3A	Well	SS-3	10.75	Split spoon	2/27/82
N-3A	Well	SS-5	20.75	Split spoon	2/27/82
N-3A	Well	SS-7	30.75	Split spoon	2/27/82
N-3A	Well	SS-9	40.75	Split spoon	2/27/82
N-3A	Well	SS-11	50.75	Split spoon	2/27/82
N-3A	Well	SS-13	60.75	Split spoon	2/27/82
N-3A	Well	SS-15	70.75	Split spoon	2/27/82
N-3A	Well	SS-17	80.75	Split spoon	2/27/82
N-3A	Well	SS-19	90.75	Split spoon	2/27/82
N-3A	Well	SS-22	100.15	Split spoon	2/27/82
N-3A	Well	SS-30	138.75	Split spoon	2/27/82
S-2	Well	SS-2	5.5	Split spoon	2/19/82
S-2	Well	SS-6A	25.3	Split spoon	2/19/82
S-2	Well	SS-10	45.75	Split spoon	2/19/82
S-8	Well	SS-4	15.7	Split spoon	3/02/82
S-8	Well	SS-9	40.75	Split spoon	3/02/82
S-8	Well	SS-12	55.75	Split spoon	3/02/82
S-8	Well	SS-14	65.55	Split spoon	3/02/82
S-8	Well	SS-17	80.55	Split spoon	3/02/82
S-11	Bore	CS-1	17.45	Split spoon	2/22/82
S-11	Bore	CS-2	36.95	Split spoon	2/22/82
S-11	Bore	SS-16	76.25	Split spoon	2/22/82
S-1	Well	SS-2	5.75	Split spoon	5/27/82
S-1	Well	SS-3	10.6	Split spoon	5/27/82
S-1	Well	SS-4	15.6	Split spoon	5/27/82
S-1	Well	SS-5B	20.7	Split spoon	5/27/82

stream-deposited. Three of the proposed sampling sites in the North Area (the dry wash entering the south boundary near N-7, South Willow Creek, and the mouth of the unnamed wash near N-8) were eliminated because there was no evidence of recent sediment deposition. These sites were replaced by taking sediment samples from the industrial waste pond (N-SD3), and the radioactive waste storage area (N-SD4).

Sediment samples were taken with a stainless-steel spade, which was cleaned carefully with paper towels and then rinsed thoroughly with distilled water. Samples were taken from a depth of approximately 3 to 5 inches below ground surface. Samples were placed in wide mouth amber glass jars with Teflon-lined plastic lids, were stored at 4°C, and were sent to the laboratory within 48 hours of collection. Sampling locations were marked with 4-foot wooden stakes painted fluorescent orange.

Table 2 is a list of sediment samples taken for chemical analysis.

B-2.3 Surface-Water Samples

Six surface water samples, three from the North Area and three from the South Area were collected for chemical analysis. The samples were all from standing water or depot effluents; no flowing streams were found at this time.

Samples were collected by partially immersing the containers in standing or flowing water. The types of containers and preservatives used are listed in Section B-4. Care was taken to prevent loss of preservative.

Field temperature, pH, Eh, and specific conductance were measured for all surface water samples. Because most samples consisted of standing water from ponds or lagoons, or the flow was very low, these measurements were not taken

Table 2. Sediment Samples for Chemical Analysis

Site I.D.	Site Type	Field Sample No.	Sample Depth (ft)	Sample Method	Date Collected
N-SD1	Ditch	N-SD1	0.5	Grab	4/3/82
N-SD2	Ditch	N-SD2	0.5	Grab	4/3/82
N-SD3	Pond	N-SD3	0.5	Grab	4/4/82
N-SD4	Ditch	N-SD4	0.5	Grab	5/26/82
S-SD1	Ditch	S-SD1	0.5	Grab	4/4/82
S-SD2	Ditch	S-SD2	0.5	Grab	4/4/82
S-SD3	Ditch	S-SD3	0.5	Grab	4/4/82
S-SD4	Ditch	S-SD4	0.5	Grab	4/4/82
N-1	Surface	GS-1	0.75	Grab	4/4/82

in situ. Instead, a beaker was rinsed out several times with a sample of surface water, and then a portion was obtained to thermally equilibrate the buffer solutions. A fresh sample was then obtained for the actual field measurement.

Samples were stored at 4°C immediately, and delivered to the laboratory within 48 hours of collection. Table 3 is a list of the surface water samples collected for analysis.

B-3 Methodology for the Collection of Ground-water Samples

B-3.1 General

Ground-water samples were collected using one of the following techniques: 1) bailing the well using a stainless steel and teflon bailer; 2) pumping the well using a nitrogen-driven double-acting piston pump made of stainless steel and Teflon (the Bennett pump); or 3) pumping the well by means existing pumps in water supply wells. This latter case occurred for the seven existing wells that we sampled on the Depot. We had no control over pump or casing materials, or over drilling and construction methods. Therefore, a lower degree of confidence is placed on the analytical results obtained for the existing wells than on those obtained for newly constructed monitoring wells.

Ensuring that representative aquifer samples could be obtained is an extremely important part of the ground-water sampling procedure. We employed several approaches in determining whether sufficient water had been evacuated from the well to produce a proper sample. Generally, procedures developed by Ertec for USATHAMA for the Rocky Mountain Arsenal sampling program were followed. In the South Area this was not a critical problem. All wells were pumped dry repeatedly during development because the transmissivity of the aquifer zones

Table 3. Surface-Water Samples for Chemical Analysis

Site I.D.	Site Type	Field Sample No.	Sample Depth (Feet)	Sample Method	Date Collected
N-SW1	Pond	N-SW1 N-SW1R	0.5 0.5	Grab Grab	4/5/82 6/23/82
N-SW2	Pond	N-SW2	0.5	Grab	4/14/82
N-SW3	Lagoon	N-SW3	0.5	Grab	6/23/82
S-SW1	Pond	S-SW1 S-SW1R	0.5 0.5	Grab Grab	4/6/82 6/24/82
S-SW2	Standing Water	S-SW2	0.5	Grab	4/15/82
S-SW3	Ditch	S-SW3	0.5	Grab	5/4/82

penetrated by the wells in this area is rather low. Since drilling fluids were not used during drilling or construction of any of these wells, pumping the wells dry guarantees recovery of 100 percent aquifer water. However, since these wells, for the most part, were not sampled within one day after being pumped dry, there is a possibility that some of the more highly volatile contaminants might have been lost. Therefore, wells that were not pumped dry within 24 hours of being sampled were closely monitored using flow-through cell procedures. This has been the case for wells in the North Area and all existing wells. The flow-through cell, discussed at length in the technical plan, allows for the measurement of pH, Eh (oxidation-reduction potential), EC (electrical conductivity), and temperature without allowing exposure to oxygen in the air and the subsequent chemical changes that immediately would occur. These parameters were monitored carefully while water was being evacuated from each well. When readings stabilized sufficiently to ensure that pumped water was representative aquifer water, samples were collected. Immediately following sample collection, parameters were again monitored to note any changes that might have occurred. If significant changes occurred in any of the parameters, the well was resampled when the parameters again stabilized. Table 1 lists the wells sampled along with method and date of sampling and amount of water evacuated before the sample was collected.

Ground-water samples at each site were collected in a series of nine sampling containers supplied by UBTL. These containers were pre-packaged with preservatives added by the lab. The type of container and preservative added for specific analytes are included in Appendix B-4. The only filtering done in the field was for the sample used for metals analyses. Filtering was accomplished using a Geofilter filter apparatus with 0.45 m filters. All samples were

Table 1. Ground-water Sampling

Volume of Water Evacuated Well No. No. Casing Volumes Gallons Date Sampled How Sampled NORTH AREA Well 1 >10* >1000 3/30/82 existing pump Well 2 >10* >1000 3/30/82 existing pump Well 4 >10 2750 3/31/82 existing pump Well 5 16 1500 4/14/82 existing pump Well 6 >10* > 250 4/4/82 existing pump N-2C dry 24 6/24/82 Bennett pump N-3A0.5 36 4/5/82 Bennett pump N-3Bdry 2.6 6/23/82 Bailed N-42.2 40 6/24/82 Bennett pump N-8B (1) 0.9 39 5/3/82 Bennett pump N-8B (2) 0.7 38 5/3/82 Bennett pump SOUTH AREA Well 1 >10* >1000 3/31/82 existing pump Well 3 >16 8540 4/15/82 existing pump S-1 2.5 27 6/24/82 Bennett Pump S-2 1.3 22 4/29/82 Bennett Pump S-3 3.2 44 4/29/82 Bennett Pump S-4 dry 21 5/2/82 Bailed S-5 1 16 5/2/82 Bennett Pump S-6 dry 76 4/28/82 Bennett Pump S-7 1 21 5/3/82 Bennett Pump S-8 dry 8.6 5/3/82 Bennett Pump S-10 1 15.4 5/2/82 Bennett Pump S-12 dry 13.2 4/28/82 Bennett Pump S-14 3 51 5/2/82 Bennett Pump

^{*} pumped regularly for Depot use

collected using teflon tubing. The sampling procedure was amended slightly during the sampling program to improve sample quality for the metals samples. Instead of supplying preservative spiked sample containers for this sample, the laboratory supplied reagent grade concentrated nitric acid which was added to the sample after collection. This prevented leaching of nickel and zinc from the sample bottles. Such leaching was found to cause significant nickel and zinc in the blank samples analyzed during the qualitative procedures used by the laboratory.

At the time of sample collection, blanks and duplicates also were included for QC purposes. A duplicate sample was taken when sampling existing Well No. 2 in the North Area. As part of the field quality control, a blank, consisting of distilled water and preservatives supplied by the laboratory, accompanied the sampling of existing well USGS No. 2. Upon sampling for each analyte, this blank container would be opened while sampling and resealed immediately after sampling was completed. The blank and duplicate were sent to the lab and analyzed along with the collected samples.

B-3.2 Existing Wells

Of the six existing wells in the North Area, the five with pumps were sampled. Well No. 3, located in the Maintenance and Supply Area, did not have a pump. This well was not sampled for two reasons: 1) it is located centrally between Wells 1 and 2, and is close enough to these two wells to be considered superfluous; and 2) because of its large diameter, it contains too much water to evacuate with a small volume pump before a representative aguifer sample could be collected.

Of the three existing wells in the South Area, only two were sampled. Well No. 2 was not sampled because of its close proximity to Well No. 1.

Sampling methodology for these existing wells consisted of:

- a. Obtaining the proper fittings to tap into the discharge line ahead of the chlorinator.
- b. Running the discharge through teflon tubing into the flow-through cell to monitor pH, Eh, EC, and temperature properly.
- c. Pump the well sufficiently long to evacuate the required amount of water.
- d. Collect the sample in proper sampling container; preserve, filter, store, and ship to lab.
- e. Proper cleaning of tubing, filter apparatus, and fittings.

B-3.3 Wells Sampled by Bailer

Because of very slow recharge and high suspended solids content, a stainless steel and teflon bailer was used to collect ground-water samples from Wells S-4 and N-3B. The procedure for sampling these wells was modified considerably since continuous monitoring could not be accomplished. After thoroughly cleaning the bailer with soapy water, acetone, and repeated distilled water rinses, we carefully lowered it into the well with a stainless steel wire. The bailer then was drawn out of the well and the water poured into a container to be disposed of after sampling was completed. This was repeated until enough water was removed from the well to ensure a minimum loss of volatile constituents from the water remaining to be sampled. Eh, pH, EC, and temperature were monitored periodically from the bailer samples. After a sufficient volume of water was evacuated from the well, samples were collected

by carefully decanting the water into the sample containers. Special care was taken when collecting the sample for volatile organics in that the sample was collected after decanting the initial third of the water from the bailer and only using the interior third. The metals sample was filtered directly from the bailer using new teflon tubing for each sample. Three bailer volumes were required to obtain the required volume for the metals samples.

B-3.4 Wells Sampled Using the Bennett Pump

The Bennett pump is an all stainless steel and teflon positive displacement pump using nitrogen to drive a double-acting piston which delivers a steady stream of water at the rate of 1/2 gallon per minute from depths as great as 625 feet. The only materials that contact water are stainless steel and Teflon. This pump was used in all monitoring wells except those bailed as discussed in the previous section. After thorough cleaning with soapy water, acetone, and repeated distilled water rinses, we carefully lowered the pump down the well until the water level indicator sounded. The well was then pumped from approximately 2 feet below the surface until the pH, Eh, EC, and temperature stabilized. The pump was lowered gradually as the water level dropped. At this time the samples were taken. Eh, pH, EC, and temperature were again measured after samples had been obtained. If significant differences in any of the readings had occurred, the well would be resampled after stabilization of Eh, pH, EC, and temperature. After use the pump was cleaned by the same procedure used prior to sampling.

This sampling procedure was modified slightly for two wells in the North Area, well N-3A and Well N-8B. In these wells the pump was lowered to a position in the center of each screened interval. The well was then pumped until monitoring parameters stabilized and a sample then was taken. From Well N-8B two

samples were taken using this procedure because there are two sets of screens that are open to two separate aquifer zones.

B-4 Methodology for Chemical Analyses

All laboratory analyses were performed by Utah Biomedical and Testing

Laboratory (UBTL) of Salt Lake City, Utah, except for radiological analyses

which were performed by Controls for Environmental Pollution (CEP), Los

Alamos, New Mexico, under supervision and control of UBTL. This section summarizes the laboratory protocols, analytical methods, and quality control used

by UBTL for the Tooele Army Depot Survey. Table 1 is a project summary, Table

2 summarizes analyses by lot, and Table 3 is a summary of lots by analysis.

Table 1. Project Summary

Project Title

Chemical Analysis for Tooele Army Depot
Contract No. DAAG 49-81-C-0192

UBTL Project Director

Sim D. Lessley, Ph.D.

					~		
Progress Summary: Samples	Jan	Feb	Mar	Apr	May	Jun	Jul
Number of Certification Samples Analyzed*	56	16	8	0	4	0	0
Number of Water Samples Received	0	0	3	15	9	6	0
Number of Soil Samples Received	0	6	17	8	0	5	0
Number of Soil Leach Tests Performed	0	6	17	8	0	5	С
Number of Soil Leach Samples Generated	0	48	70	64	0	33	0
Number of Field Samples Analyzed by Method:							
Volatiles	0	0	0	10	13	0	7
Semi Volatiles	0	0	12	6	0	27	10
Explosives	0	0	6	25	25	4	5
NG and PETN	0	0	23	8	25	9	0
Oil and Grease	0	0	Ú	0	25	0	5
Metals - ICP	0	0	6	0	6	20	22
Metals - GF/AA	0	0	12	20	13	0	9
Mercury	0	0	12	20	13	6	4
Sodium	0	0	6	26	13	0	9
Anions	0	3	9	24	9	9	0
Cyanide	0	0	0	0	25	0	5
Gross Alpha and Beta	0	_0	0	0	0	47	10
Total Number of Field Samples Analyzed	0	3	86	139	167	122	86
Number of QC Spikes Analyzed	0	1	16	37	20	19	27
Number of QC Blanks Analyzed	0	_2	25	_33	16	22	18
Total Number of Samples Analyzed (Field & QC)	0	6	127	209	203	163	131

^{*}Note eighteen additional recertification samples were analyzed in September 1982.

Table 2.
Summary of Analyses by Lot

Lot	Matrix	Analysis	Date Analyzed	Field Samples	QC Samples	Blanks	Total Samples
AAA	Leach	Anions	02/26/82	3	1	1	5
AAB	Leach	Anions	03/01/82	3	1	1	5
AAC	Leach	NG & PETN	03/05/82	6	1	2	9
AAD	Leach	5 Explosives	03/10/82	6	1	2	9
AAE	Leach	Semi Volatiles	03/12/82	6	1	2	9
AAF	Leach	Mercury	03/05/82	6	1	2	9
AAG	Leach	Metals-ICP	03/11/82	6	2	2	10
AAH	Leach	Anions	03/05/82	5	1	1	7
AAI	Leach	Metals GF/AA	03/16/82	6	1	2	9
AAJ	Leach	Sodium	03/10/82	6	1	2	9
AAK	Leach	Mercury	03/11/82	6	1	2	9
AAL	Leach	Anions	03/11/82	1	1	1	3
AAM	Leach	NG & PETN	03/17/82	11	1	1	13
AAN	Leach	NG & PETN	03/17/82	6	1	1	8
AAO	Leach	5 Explosives	04/01/82	11	1	1	13
AAP	Leach	5 Explosives	04/01/82	6	1	1	8
AAQ	Leach	Semi Volatiles	03/18/82	6	1	2	9
AAR	Leach	Sodium	04/07/82	6	1	2	9
AAS	Leach	Metals GF/AA	03/25/82	6	1	2	9
AAT	Leach	Metals ICP	05/13/82	6	2	2	10
AAU	Water	Anions	04/03/82	4	2	1	7
AAV	Water	Anions	04/06/82	2	1	1	4
AAW	Water	Volatiles	04/06/82	4	7	0	11
AAX	Water	Volatiles	04/06/82	2	3	2	7
AAY	Water	Mercury	04/07/82	6	2	1	9
AAZ	Water	Semi Volatiles	04/29/82	6	2	1	9

Lot	Matrix	Analysis	Date Analyzed	Field Samples	QC Samples	Blanks	Total Samples
ABA	Water	Anions	04/07/82	2	1	0	3
ABB	Water	NG & PETN	04/13/82	8	2	0	10
ABC	Water	5 Explosives	04/16/82	8	2	0	10
ABD	Water/Leach	Anions	04/16/82	12	2	4	18
ABE	Water	Mercury	04/20/82	14	2	6	22
ABF	Water	Volatiles	04/21/82	4	1	1	6
ABG	Water/Leach	NG & PETN	05/05/82	12	2	4	18
ABH	Water/Leach	Explosives	05/10/82	12	2	4	18
ABI	Water/Leach	Semi Volatiles	06/08/82	14	2	4	20
ABJ	Water/Leach	Metals GF/AA	04/28/82	20	3	5	28
ABK	Water	Sodium	04/28/82	20	3	7	3 0
ABL	Water/Leach	Metals ICP	06/03/82	20	4	5	29
ABM	Water	Cyanide	05/05/82	12	3	2	17
ABN	Water	Oil & Grease	05/06/82	12	2	2	16
ABO	Water	Anions	04/30/82	4	1	0	5
ABP	Water	Anions	05/04/82	4	1	0	5
ABQ	Water	Anions	05/06/82	5	1	0	6
ABR	Water	Cyanide	05/11/82	13	1	0	14
ABS	Water	Mercury	05/07/82	13	1	0	14
ABT	Water	Semi Volatiles	06/24/82	13	1	0	14
ABU	Water	Volatiles	05/06/82	13	1	0	14
ABV	Water	5 Explosives	05/22/82	13	1	0	14
ABW	Water	NG & PETN	05/27/82	13	1	0	14
ABX	Water	Oil & Grease	05/11/82	13	0	0	13
ABY	Water	Sodium	05/12/82	13	1	1	15
ABZ	Water	Metals GF/AA	05/12/82	13	1	1	15

Lot	Matrix	Analysis	Date Analyzed	Field Samples	QC Samples	Blanks	Total Samples
ACA	Water	Metals ICP	07/27/82	13	6	1	20
ACB	Leach	Anions	06/05/82	4	1	1	6
ACC	Leach	Semi Volatiles	07/14/82	4	1	1	6
ACD	Leach	5 Explosives	06/15/82	4	1	1	6
ACE	Leach	NG/PETN	06/17/82	4	1	1	6
ACF	Leach	Mercury	07/15/82	4	1	1	6
ACG	Water	Anions	06/25/82	5	2	1	8
ACH	Water	NG & PETN	06/29/82	5	2	1	8
ACI	Water	5 Explosives	07/01/82	5	2	1	8
ACJ	Water/Leach	Metals ICP	07/27/82	9	3	2	14
ACK	Water/Leach	Mercury	06/30/82	6	2	1	9
ACL	Water	Volatiles	07/02/82	7	2	1	10
ACM	Water	Cyanide	07/01/82	5	2	1	8
ACN	Water	Semi Volatiles	07/14/82	6	2	1	9
ACO	Water	Oil & Grease	07/01/82	5	1	1	7
ACP	Water/Leach	Metals GF/AA	07/07/82	9	2	2	13
ACQ	Water/Leach	Sodium	07/02/82	9	2	2	13
ACR	Water/Leach	Gross Alpha & Beta	06/18/82	47	3	7	57
ACS	Leach/Water	Gross Alpha & Beta	07/22/82	10	3	4	17

Table 3. Summary of Lots by Analysis

Analysis

Lots

Volatiles (2J)	AAW,	AAX,	ABF,	ABU,	ACL				
Semi Volatiles (3W)	AAE,	AAQ,	AAZ,	ABI,	ABT,	ACC,	ACN		
5 Explosives (2B)	AAD,	AAO,	AAP,	ABC,	ABH,	ABV,	ACD,	ACI	
NG & PETN (6B)	AAC,	AAM,	AAN,	ABB,	ABG,	ABW,	ACE,	ACH	
Metals ICP (3T)	AAG,	AAT,	ABL,	ACA,	ACJ				
Metals GF/AA (7T)	AAI,	AAS,	ABJ,	ABZ,	ACP				
Mercury CV/AA (1D)	AAF,	AAK,	AAY,	ABE,	ABS,	ACF,	ACK		
Sodium (1M)	AAJ,	AAR,	ABK,	ABY,	ACQ				
Anions (2P)		AAB, ABQ,			AAU,	AAV,	ABA,	ABD,	ABO,
Cyanide (4K)	ABM,	ABR,	ACM						
Oil & Grease (00)	ABN,	ABX,	ACO						
Gross Alpha & Beta (30)	ACR,	ACS							

This section contains the laboratory protocols used for the handling of samples and data from the Tooele Army Depot Survey. They are presented in the following order:

Sample Logging
Sample Lotting
Soil/Sediment Quartering Procedure
Soil Sample Handling And Wetting Volume Determination
Response to Out-Of-Control QC Data
Water Sample Containers
Soil Leach Sample Containers

Sample Logging

Soil Sample Log In

The soil samples will be received in styrofoam boxes packed with blue ice. As each soil sample is received it will be inspected and the following information entered into the first section of the sample logbook.

- 1. Date of arrival of sample at the laboratory.
- 2. The field identification numbers of the sample.
- Sample matrix (soil, sediment).
- 4. Sampling date.
- 5. Date leaching procedure performed.
- 6. Analytical procedure to be run.
- 7. Observations concerning sample conditions (e.g. broken containers, leakage, lack of temperature control, etc.).

The soil samples will be stored in the walk-in refrigerator pending analysis. The blue ice must be removed before the samples are stored in the refrigerator.

Water Sample Log In

Each water sample will consist of several glass and plastic bottles packed in a styrofoam box with blue ice. Before storage of samples in the refrigerator the blue ice must be removed from the box. As each water sample is received it will be inspected and the following information entered into the first section of the sample logbook:

- 1. Date of arrival of sample at the laboratory.
- 2. The field identification numbers of the sample.
- Sample matrix (surface water, ground water).
- 4. Sampling date.
- 5. Date split made.
- 6. Analytical procedure to be run.
- 7. Observation concerning sample conditions (e.g. broken containers, leakage, lack of temperature control, etc.).

In cases where samples are split (see below) the field identification number will be attached to each split container. The water samples will be stored in the walk-in refrigerator or in the sample room, as appropriate, grouped according to analytical procedure.

Water samples for explosives and metals will be split for analysis.

The water sample for explosives will be collected in a 1L amber glass bottle. The split will be made by pouring 250 mL directly from the 1L bottle into a 250 mL amber glass bottle. The sample remaining in the 1L bottle (approximately 750 mL) is to be analyzed for the five explosives: TNT, 2,4-DNT, 2,6-DNT, RDX and Tetryl. The sample in the 250 mL bottle is to be analyzed for NG and PETN. A preservative of 2-nitrodiphenylamine in isopropanol is added to each NG/PETN sample, resulting in a final concentration of 0.05 mM for the 2-nitrodiphenylamine.

The water sample for metals will be collected in a 1L plastic bottle. The splits will be made by pouring directly from the 1L bottle. The splits are as follows:

Split Container	Volume	Analytical Procedure		
30 mL Plastic	30 mL	Metals-GF/AA		
250 mL Plastic	250 mL	Mercury		
30 mL Plastic	30 mL	Sodium		
lL Plastic	remaining volume	Metals-ICP		

Sample Lotting

Lot size is limited by storage time or the number of samples that can be processed through the method in one day.

When a particular lot is ready to analyze, the samples are removed from the storage area. The QC's are prepared in the same type of sample containers and randomly distributed among the samples. The three-letter, three-digit codes are tagged to the sample. The three letter, three-digit code for each sample is written on the field tag before the field tag is placed in the lot file folder. The following information is recorded for each sample in the second section of the logbook: Laboratory Number, Field (or QC) Number, Matrix, Analytical Procedure, Chemist, Date Analyzed.

Soil/Sediment Quartering Procedure

Each soil sample will be sieved through a 9.5 mm stainless steel sieve, if necessary. The sample will then be poured out on a double thickness of heavy duty aluminum foil (dull side up) and mixed with a stainless steel spatula. The sample will be quartered and opposite quarters returned to the sample bottle. The remaining pair of opposite quarters are combined, mixed and again quartered. Opposite quarters are returned to the sample bottle. This process is continued until approximately 200 g of soil is left. The sample is quartered and opposite quarters are combined, weighed and taken for extraction. The remaining pair of opposite quarters are combined, weighed and taken for the determination of wetting volume.

Each sediment sample will be poured through the 9.5 mm stainless steel sieve, if necessary. The sample will be stirred very well and an

appropriate amount taken for leaching. The wetting volume is assumed to be zero. Therefore, no wetting volume determination will be performed for sediment.

Soil Sample Handling And Wetting Volume Determination

Break down soil lumps. Do not grind, just break down to natural size particles. Screen through the 9.5 mm screen. Weigh and discard what was retained on the sieve. Weigh what went through the sieve.

Perform the necessary splits to obtain approximately 100 g for extraction and approximately 100 g for determination of wetting volume. In the event that a very small sample is received as little as 25 to 50 grams can be used to determine the wetting volume.

The wetting volume is determined as follows:

- Weigh approximately 100 g of soil into a 250 mL plastic centrifuge bottle. Add a measured volume of water - 100 mL for 100 g.
- 2. Cap and shake well. Allow to stand for at least one hour.
- 3. Centrifuge.
- 4. Decant the supernatant water and measure its volume with a graduated cylinder.
- 5. Determine the wetting volume by subtraction:
 Wetting Volume = (Original Volume of Water) (Volume of Supernatant)

The wetting volume is used to determine the total leaching volume.

Response To Out-of-Control QC Data

All analytical data should be reduced as soon as possible and routed to the quality assurance specialist (QAS).

In the event that the data is out of control, analyses will be discontinued until the analytical system is demonstrated to be under control. All data obtained since analysis of the latest in-control QC samples will be considered invalid. Efforts will be directed toward the

determination of the cause of the problem(s) and required corrective actions will be implemented to reinstate a reliable analytical system. Once problems are identified and corrected, data considered to be invalid will be evaluated. If identified errors can be corrected without the reanalysis of the samples (as, for example, in the case of calculation errors or improper processing of reliable raw analytical data), corrections will be effected without additional analytical work. All samples for which reanalysis is the only reasonable corrective measure, will be reanalyzed by UBTL if sufficient sample is available and if the integrity of the remaining portion of the sample is intact. Resampling and reanalysis will be required to correct out-of-control results if the conditions noted in the preceding cannot be satisfied.

As noted, the UBTL QAS will carefully monitor (on a real-time basis) the results of the control samples employed for this contract work, and in the event that a QC result is out-of-control, specific steps will be implemented to identify and correct the deficiency. The steps, to be pursued jointly by the analyst and the QAS, are as follows:

- 1. Check all data processing procedures and calculations.
- Check blank samples for identification of possible interferences or other problems.
- 3. Check instrumentation performance (if applicable) by observing the response of the instrument while processing a sample material for which the expected response is known. Operating conditions must be similar to those used for analysis of the samples under consideration.
- 4. Check the original standard preparation procedures by preparing new standards, obtaining a new standard calibration curve from the new data, and comparing the new standard curve with the original standard calibration curve.
- 5. Check the integrity of the original QC samples by preparing new QC samples following the same procedures, and analyzing the new QC samples.

- 6. Carefully review raw data (e.g., recorder output, chromatograms, computer output) in an effort to identify interferences, unusual signals (unusual peak shapes, etc), or other factors which could produce inaccuracies.
- 7. Reanalyze the samples with new standards and new QC samples if sufficient field sample material is available. The entire analytical process including, for example, extractions, digestions, etc. should be repeated.
- 8. Any other special procedures required to address recalcitrant problems not solved by the preceding steps.

Water Sample Containers

Table 4 lists the containers to be used in the collection and analysis of water samples from the Tooele Army Depot Survey.

Table 4. Water Sample Containers

Methods -	Sampling	Container	Split	Container	Volume	Volume
Methods -	Size	Material	Size	Material	to Analyst	for Analysis
Volatiles (624)	2 x 40 mL	Amber Glass	-	-	2 x 40 mL	2 x 5 mL
Semi-Volatiles (625)	1L	Amber Glass	-	-	1L	1L
Explosives	1L	Amber Glass	lL	Amber Glass	750 mL	500 mL
NG & PETN	TL	Amber Glass	250 mL	Amber Glass	250 mL	100 mL
Oil & Grease	1L	Amber Glass	-	-	1L	1L
Metals-ICP			1L	Plastic	690 mL	250 mL
Metals - GF/AA	1L	Plastic	30 mL	Plastic	30 mL	20 mL
Mercury - CV/AA	115	Plastic	250 mL	Plastic	250 mL	100 mL
Sodium - AA			30 mL	Plastic	30 mL	20 mL
Anions - IC	30 mL	Plastic	-	-	30 mL	5 mL
Cyanide	1L	Plastic	-	-	1L	500 mL
Gross Alpha & Beta	500 mL	Plastic	-	-	500 mL	250 mL

Soil Leach Sample Containers

Table 5 lists the containers to be used in the collection and analysis of soil leach samples from the Tooele Army Depot Survey.

Table 5.
Soil Leach Sample Containers

Method -	Leach	Container	Volume	Volume	
	Size	Material	to Analyst	for Analysis	
Volatiles (624)	-	-	_	-	
Semi-Volatiles (625)	250 mL	Amber Glass	200 mL	200 mL	
Explosives	250 mL	Amber Glass	100 mL	100 mL	
NG & PETN	40 mL	Amber Glass	40 mL	20 mL	
Oil & Grease	-	-	-	-	
Metals-ICP	250 mL	Plastic	50 mL	50 mL	
Metals - GF/AA	30 mL	Plastic	30 mL	20 mL	
Mercury - CV/AA	30 mL	Plastic	30 mL	20 mL	
Sodium - AA	30 mL	Plastic	30 mL	20 mL	
Anions - IC	30 mL	Plastic	30 mL	20 mL	
Cyanide	-	-	-	-	
Gross Alpha & Beta	250 mL	Plastic	125 mL	50 mL	

Remainder is stored as follows: 1) One amber glass 250 mL bottle filled to top; 2) The rest in a 250 mL plastic bottle, preserved with nitric acid.

Qualitative Analysis Methods

For the screening part of the Tooele Army Depot Survey, twelve laboratory analyses were performed. Eleven analyses were certified at the qualitative level. The analysis for oil and grease was performed without certification. Table 6 summarizes the results of the qualitative analysis certifications. The soil leach samples were analyzed by the same method used for water samples with adjustments made for sample size and detection limit as suggested in the Solid Waste Leaching Procedure (SWLP).

Semi-Quantitative Analysis Methods

At the request of Ertec Western, Inc., UBTL proceeded with the recertification of the analytical methods at the semi-quantitative level. It was the intent of the Project Officer, Mr. Donald Campbell, that no resampling be performed and that the semi-quantitative recertification be applied to the data already reported by the laboratory.

UBTL reviewed the analytical data and recommended that the methods for NG & PETN (6B) and mercury (1D) not be recertified because none of the field samples contained those analytes at or above the detection limit.

When the qualitative analysis certifications were being performed, the UBTL chemists were directed by the Chemical Analysis Program Manager, Dr. Sim D. Lessley, to analyze six samples at the levels required for the semi-quantitative recertification, if time permitted. This was done with a view to reducing recertification costs for the resampling portion of the program. Upon careful review of the original certification data, it was discovered that the UBTL chemists had been able to run the six levels necessary for the semi-quantitative certification of several methods while the qualitative analyses were being certified. These methods were:

Method 2J (Volatiles in water by HP-5985B GC/MS) Method 7T (Zinc, Arsenic and Nickel in water by GF/AA) Method 1M (Sodium in water by AA)

Table 6. Certification of Qualitative Analyses

Method	Method #	Analyte	Analyte Code	ug/L LOD Water	µg/I LOD Soil Le		Slope ecovery)
						JMR	RWW
Volatiles	2 J	Benzene	C6H6	1	-	1.27	1.22
(624)		Bromomethane	CH3BR	1	_	1.16	0.367
		Chlorobenzene	CLC6H5	1	_	1.03	1.39
		1,2-Dichloroethane	12DCLE	1	-	1.24	0.767
		Trans-1,2-Dichloroethene	T12DCE	1	-	1.24	0.903
		1,1,2,2-Tetrachloroethane	TCLEA	1		1.42	1.01
		1,1,1-Trichloroethane	111TCE	1	-	1.41	0.931
		Trichloroethene	TRCLE	1	_	1.28	1.45
Semi-Volatiles	3W	Hexachloroethane	CL6ET	20	100	JMR	RWW
July Voluciacy	24	Naphthalene				0.333	0.592
		Nitrobenzene	NAP	2	10	0.530	0.76
<u> </u>			NB	8	40	0.516	0.80
		3,5-Dinitroaniline	35DNA	20	100	1.19	0.559
		2-Amino-4,6-DNT Fluoranthene	2A46DT	20	100	1.38	1.46
			FANT	2	10	1.01	1.00
		3-Nitrotoluene	3NT	10	50	0.792	1.00
		Diethylphthalate	DEP	2	10	1.04	0.99
		Alpha-BHC	ABHC	20	100	0.983	1.00
		p,p'-DDT	PPDDT	2	10	1.07	1.00
		Dieldrin	DLDRN	2	10	1.04	0.70
		Lindane	LIN	20	100	1.03	1.02
		Heptachlor	HPCL	8	40	0.998	0.76
		Aroclor-1016	PCB016	70	350	0.925	1.00
		Aroclor-1262	PCB262	100	500	0.991	0.73
		2,4-Dimethylphenol	24DMPN	20	100	0.028	0.60
		2,4-Dinitrophenol	24DNP	30	150	1.02	1.07
		2-Methyl-4,6-dinitrophenol	46DN2C	20	100	1.27	1.08
		Pentachlorophenol	PCP	20	100	1.14	1.02
		Phenol (D6)	PHEND6	3	15	0.235	0.252

Table 6. Certification of Qualitative Analyses (Continued)

Me thod	Method #	Analyte	Analyte Code	μg/L LOD Water	μg/L LOD Soil Le		lope overy)
Explosives	2B	2,4-DNT	24DNT	2	10	0.739	
		2,6-DNT	26DNT	3	15	0.741	
		2,4,6-TNT	246TNT	2	10	0.852	
		Tetryl	TETRYL	1	5	0.897	
		RDX	RDX	1	5	0.940	
NG & PETN	6B	Nitroglycerine	NG	20	100	0.643	
		PETN	PETN	5	25	1.08	
Metals-ICP	3т	Arsenic	AS	60	300	0.973	
		Beryllium	BE	0.5	3	1.06	
		Cadmium	CD	6	30	0.964	
		Chromium	CR	5	25	0.967	
		Copper	CU	6	30	1.07	
		Lead	PB	30	150	0.928	
		Nickel	NI	20	100	0.931	
		Silver	AG	8	40	0.929	
		Zinc	ZN	20	100	0.832	
Metals-GF/AA	7 T	Arsenic	AS	7	7	0.913	
		Nickel	NI	5	5	0.94	
		Zinc	ZN	1	1	1.11	
						CLM	DWB
Mercury-CV/AA	1D	Mercury	HG	0.2	1	0.915	0.909
Sodium	1M	Sodium	NA	1000	1000	0.989	

Table 6.
Certification of Qualitative Analyses (Continued)

Method	Method #	Analyte	Analyte Code	ug/L LOD Water	μg/L LOD Soil Lea		lope overy)
						LJD	REB
Anions	2P	Chloride	CL	1000	1000	0.971	1.15
		Fluoride	F	1000	1000	0.894	1.15
		Nitrate	NO3	1000	1000	1.14	1.06
		Nitrite	NO2	1000	1000	1.05	0.956
		Phosphate	P04	1000	1000	0.938	1.01
		Sulfate	S04	1000	1000	0.934	1.03
Cyanide	4K	Cyanide	CYN	10	-	0.919	
Oil & Grease	00	Oil & Grease	OILGR	5000	-	-	
Gross Alpha &	30	Gross Alpha	ALPGL	2pCi/L	10pCi/L	0.893	
Beta	30	Gross Beta	BETGL	3pCi/L	15pCi/L	0.786	

Method 2P (Anions in water by Ion Chromatography)

Method 4K (Cyanide in water by Spectrophotometry)

Method 3T (Nine metals in water by ICP)

The necessary analytical work was performed to recertify the three remaining analytical methods:

Method 3W (Semi-volatiles in water by HP 5985B GC/MS)

Method 2B (Five Explosives in water by HPLC)

Method 30 (Gross Alpha and Beta Radiation in water)

Table 7 summarizes the results of semi-quantitative analysis certifications. Note that the last column of the table contains a multiplier (certified slope divided by recertified slope) for use in converting qualitative analysis data to semi-quantitative analysis data.

The semi-quantitative analytical methods are contained in Appendix F.

0.876

Certification of Semi-Quantitative Analyses Table ,.

Method	Me thod	d Analyte	Analyte Code	Cert. LOD µg/L Water	Cert. Slope	Recert. LOD µg/L Water*	Recert. LOD µg/L Leach	Recert. Slope	Cert. Slope Recert. Slope
					(JMR)				
Volatiles	2.3	Benzene	91192	1	1.27	2	;	0.988	1.29
(624)		Bromomethane	CH3BR	1	1.16	5	1	0.943	1.23
		Chlorobenzene	СГС6Н5	-	1.03	2	;	1.08	0.954
		1,2-Dichloroethane	12DCLE		1.24	2	;	1.16	1.07
		Trans-1,2-Dichloroethene	T12DCE	1	1.24	2	}	1.03	1.20
		1,1,2,2-Tetrachloroethane	TCLEA	1	1.42	2	;	1.01	1.41
		l,1,1-Trichloroethane	111TCE	1	1.41	e	ļ	1.25	1.13
		Trichloroethene	TRCLE	7	1.28	м	1	1.18	1.08
					(11114)				
					(KWM)				
Semi-Volatiles	3W	Hexachloroethane	CL6ET	20	0.592	8(20)	100	0.791	0.748
(625)		Naphthalene	NAP	2	0.76	2	10	0.999	0.761
		Ntrobenzene	NB	8	08.0	5(8)	40	0.987	0.811
		3,5-Dinitroaniline	35DNA	20	0.559	20	100	0.842	0.664
		2-Amino-4,6-DNT	2A46DT	20	1.46	20	100	0.983	1.49
		Fluoranthene	FANT	2	1.00	3	15	0.777	1.30
		3-Nitrotoluene	3NT	10	1.00	6(10)	50	096.0	1.04
		Diethylphthalate	DEP	2	0.99	5	25	1.13	0.876

*The value given is the detection limit for which the method was certified. The value in parentheses is the detection limit to be used for the Tooele Survey.

Certification of Semi-Quantitative Analyses (Continued)

Method	Method #	d Analyte	Analyte Code	Cert. LOD ng/L Water	Cert. Slope	Recert. LOD µg/L Water*	Recert. LOD µg/L Leach	Recert. Slope	Cert. Slope Recert. Slope
		Alpha-BHC	ABIIC	20	1.00	30	150	1.12	0.893
		P, p'-DDT	PPDDT	2	1.00	2	10	0.782	1.37
		Dieldrin	DLDRN	2	0.70	4	20	1.01	0.693
		Lindane	LIN	20	1.02	30	150	0.471	2.17
		Heptachlor	ПРСГ	80	0.76	6	45	0.642	1.18
		Aroclor-1016	PCB016	70	1.00	300	1500	0.926	1.08
		Aroclor-1262	PCB262	100	0.73	300	1500	0.798	0.915
		2,4-Dimethylphenol	24DMPN	20	09.0	20	100	0.048	12.5
		2,4-Dinitrophenol	24DNP	30	1.07	40	200	0.803	1.33
		2-Methyl-4,6-dinitrophenol	46DN2C	20	1.08	40	200	0.880	1.23
		Pentachlorophenol	PCP	20	1.02	40	200	0.612	1.67
		Phenol (D6)	PHEND6	е	0.252	9	30	0.340	0.741
Explosives	2B	2,4-DNT	24DNT	2	0.739				
		2,6-DNT	26DNT	د	0.741				
		2,4,6-TNT	246TNT	7	0.852				
		Tetryl	TETRYL	1	0.897				
		RDX	RDX	-1	0.940				
NG & PETN	68	Nitroglycerine	NG	20	0.643	i	1	;	ţ
		PETN	PETN	5	1.08	1	1	1	1

The value in parentheses is the *The value given is the detection limit for which the method was certified. detection limit to be used for the Tooele Survey.

Table 7. Certification of Semi-Quantitative Analyses (Continued)

					•				
	Method		•	Cert.		Recert. LOD	Recert. LOD		Cert. Slope
Method	#	Analyte	Analyte Code	μg/L Water	Cert. Slope	µg/L Water*	μg/L Leach	Recert. Slope	Recert. Slope
Metals-ICP	3T	Arsenic	AS	09	0.973	40(60)	300	0.958	1.02
		Beryllium	36	0.5	1.06	0.3(0.5)	3	0.988	1.07
		Cadmium	9	9	0.964	9	30	0.894	1.08
		Chromium	CR	5	0.967	4(5)	25	0.927	1.04
		Copper	CU	9	1.07	9	30	0.989	1.08
		Lead	PB	30	0.928	20(30)	150	0.888	1.05
		Nickel	IN	20	0.931	7(20)	100	0.893	1.04
		Silver	AG	œ	0.929	ಹ	70	0.832	1.12
		Zinc	ZN	20	0.832	6(20)	100	0.814	1.02
Metals-GF/AA	7.1	Arsenic	AS	7	0.913	4(7)	7	0.945	9960
		Nickel	NI	5	96.0	4(5)	2	1.01	0.931
		Zinc	NZ	-	1.11		9	1.25	0.888
Mercury-CV/AA	10	Mercury	HG	0.2	0.915	0.1(0.2)	1	0.903	1.01
Sodium	M	Sodium	NA 10	1000	0.989	1000	1000	1.00	0.989
Anions	2 P	Chloride	CL 10	0001	0.971	1000	1000	0.990	0.981
		Fluoride	F 10	0001	0.894	2000	2000	1.01	0.885
		Nitrate	NO3 10	1000	1.14	2000	2000	1.03	1.11

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*The value given is the detection limit for which the method was certified. The value in parentheses is the

Table /. Certification of Semi-Quantitative Analyses (Continued)

Cert. Slope Recert. Slope	1.08	0.972	976.0	ŀ	
Recert. Slope	0.972	0.961	0.942	1	
Recert. LOD µg/L Leach	1000	2000	;	ļ	
Recert. LOD µg/L Water*	1000	2000	5(10)	1	
Cert. Slope	1.05	0.934	0.919	1	0.893
Cert. LOD Hg/L Water	1000	1000	01	2000	ALPGL 2 PC1/L BETGL 3 PC1/L
Analyte Code	N02 P04	804	CYN	OILGR	ALPGL BETGL
Analyte	Nitrite Phosphate	Sulfate	Cyanide	011 & Grease	Gross Alpha Gross Beta
Method #			4K	00	30
Method			Cyanide	011 & Grease	Gross Alpha & Beta

*The value given is the detection limit for which the method was certified. The value in parentheses is the detection limit to be used for the Tooele Survey.

QUALITY CONTROL

Quality Control Samples

Quality control samples were included in each analysis lot. The types of samples prepared by the UBTL Quality Assurance Specialist were:

Standard Water Spike - the analytes of interest were made up at the certified detection limit in standard water (standard water is defined in the USATHAMA QA/QC Manual).

Natural Water Spike - a field sample was split and the analytes of interest were made up at the detection limit in one of the splits.

Field Spike - a standard water spike was prepared, as described above, transported to the field and returned to the laboratory for analysis.

Standard Water Blank - a sample of the standard water used to prepare the standard water spikes was included as a blank.

Field Blank - a standard water blank was prepared as described above, transported to the field and returned to the laboratory for analysis.

The radiological samples represented an exception to the foregoing. In the case of gross alpha and gross beta determinations the contract laboratory, Controls for Environmental Pollution (CEP), prepared quality control samples and method blanks for each analysis lot.

Tables 8 and 9 summarize the quality control objectives for water samples and soil leach samples, respectively. In some cases there were departures from the objectives. These cases are noted individually in the lot quality control report (see below).

Table 8. Quality Control Objectives for Water Samples

Method	Std. H ₂ 0 Blank	Std. H ₂ 0 Spike	Natural Spike	Field Blank	Field	Spike
Volatiles (624)	one					
Bromomethane						
Chlorobenzene		one	none	one	one	lot
1,2-Dichloroethane						
Semi-Volatiles (625)	one					
Hexachloroethane						
3,5-Dinitroaniline		one	one	one	one	10+
Dieldrin			00	one	one	100
2,4-Dinitrophenol						
Explosives						
All analytes	one					
All analytes		one	one	one	one	lot
NG & PETN	one					
Both analytes		one	one	one	one	lot
				o ii.e	One	100
Metals - ICP	one		•			
All except silver		one	one	one	one .	lot
Silver		one	none	one	one :	lot
Metals - GF/AA	one					
All analytes	One	One	020			
, 200		one	one	one	one :	lot
Mercury - CV/AA	one	one	one	one	one :	lot
Sodium	one	one	one	one	one :	lot
Anions	one					
All analytes	One	000				_
didiyees		one	one	one	one :	lot
Cyanide	one	one	one	one	one :	lot
					· · · · ·	
Oil & Grease	one	none	none	one	non	e
Gross Alpha & Beta	0.7.5					
ooo mapma a beta	one	none	none	one	non	e

Table 9. Quality Control Objectives for Soil Leach Samples

Method	Std. H ₂ 0 Blank	Std. H ₂ O Spike
Volatiles (624) Bromomethane Chlorobenzene 1,2-Dichloroethane	none	none
Semi-Volatiles (625) Hexachloroethane 3,5-Dinitroaniline Dieldrin 2,4-Dinitrophenol	one	one
Explosives All analytes	one	one
NG & PETN Both analytes	one	one
Metals - ICP All except silver Silver	one	one one
Metals - GF/AA All analytes	one	one
Mercury - CV/AA	one	one
Sodium	one	one
Anions All analytes	one	one
Cyanide	one	one
Oil & Grease	one	none
Gross Alpha & Beta	one	none

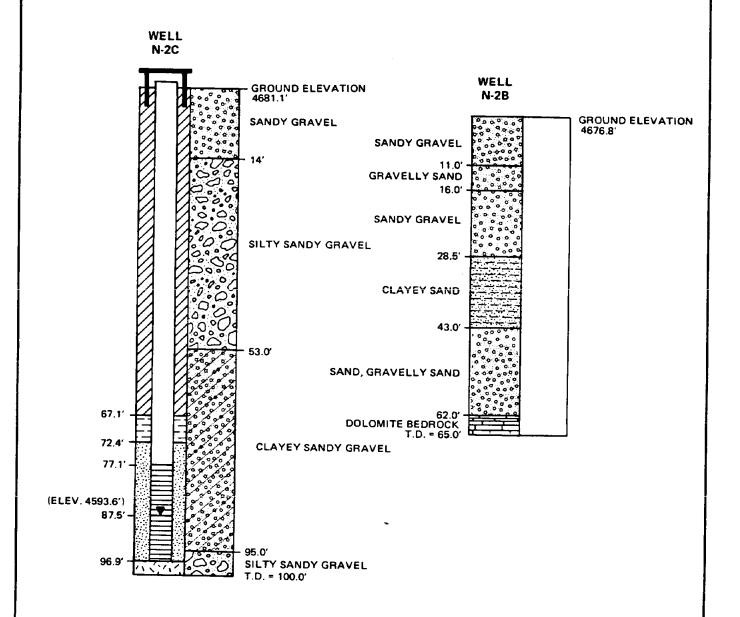
Quality Control Reports

At the request of Dr. Les Eng, Mr. Lance M. Eggenberger, (the UBTL Quality Assurance Specialist) prepared a quality control report for each lot of samples. In each report the results of the analysis of the quality control samples for the lot were presented and discussed. Copies of all of the quality control reports for the Tooele Army Depot Survey are included in Appendix G.

Quality Control Results

Also at the request of Dr. Les Eng the UBTL Quality Assurance Specialist tabulated the results of the analyses of all quality control samples by analyte. Those tabulations are presented in Appendix H. The quality control results are summarized by lot with the range of the field sample data for each lot in Appendix I.

Appendix C
Well Construction and
Lithology Summaries

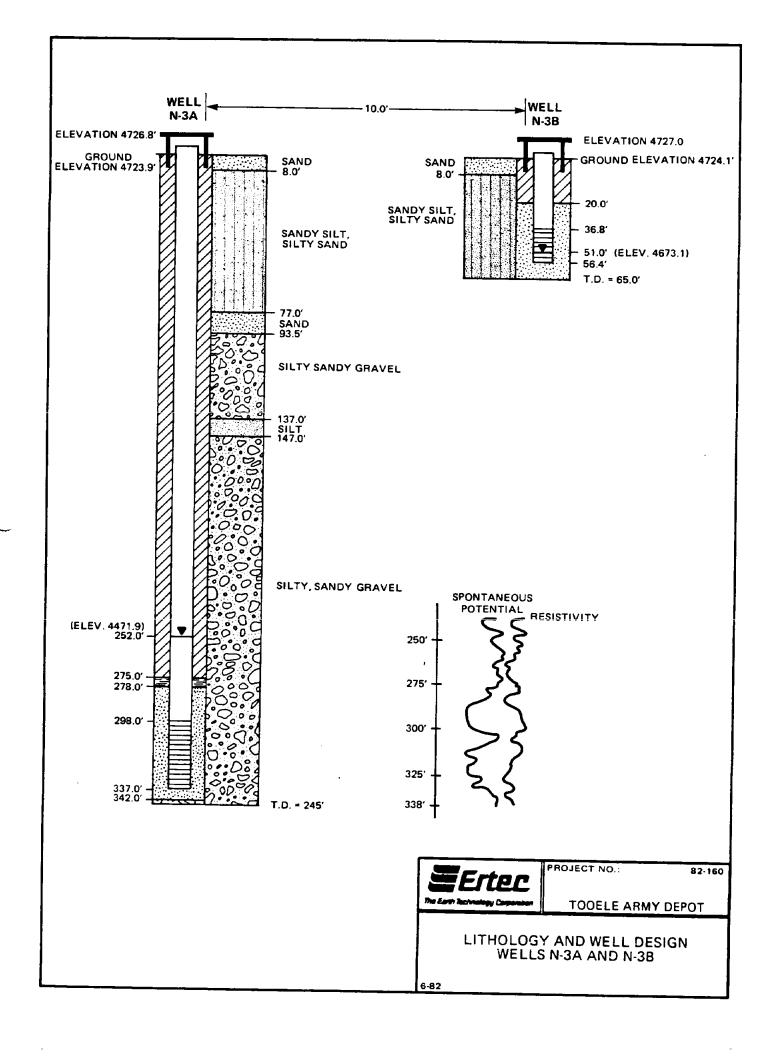


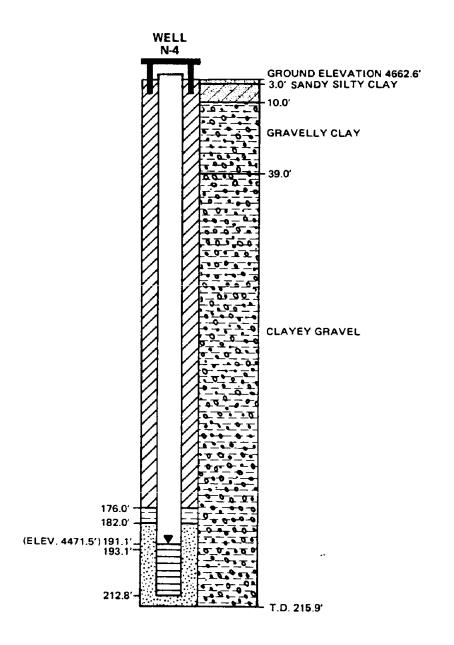


82-160

TOOELE ARMY DEPOT

LITHOLOGY AND WELL DESIGN WELLS N-2C AND N-2B



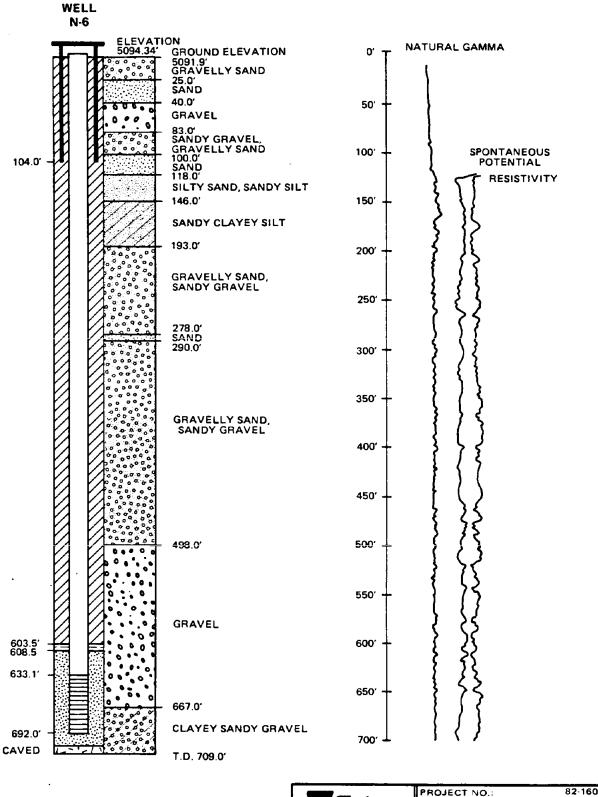




82-16

TOOELE ARMY DEPOT

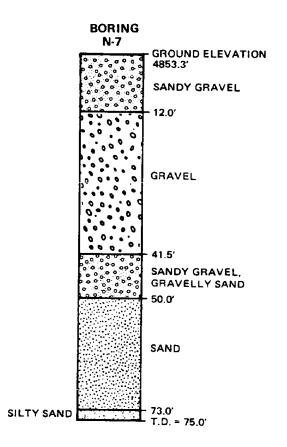
LITHOLOGY AND WELL DESIGN WELL N-4



82-160

TOOELE ARMY DEPOT

LITHOLOGY AND WELL DESIGN WELL N 6

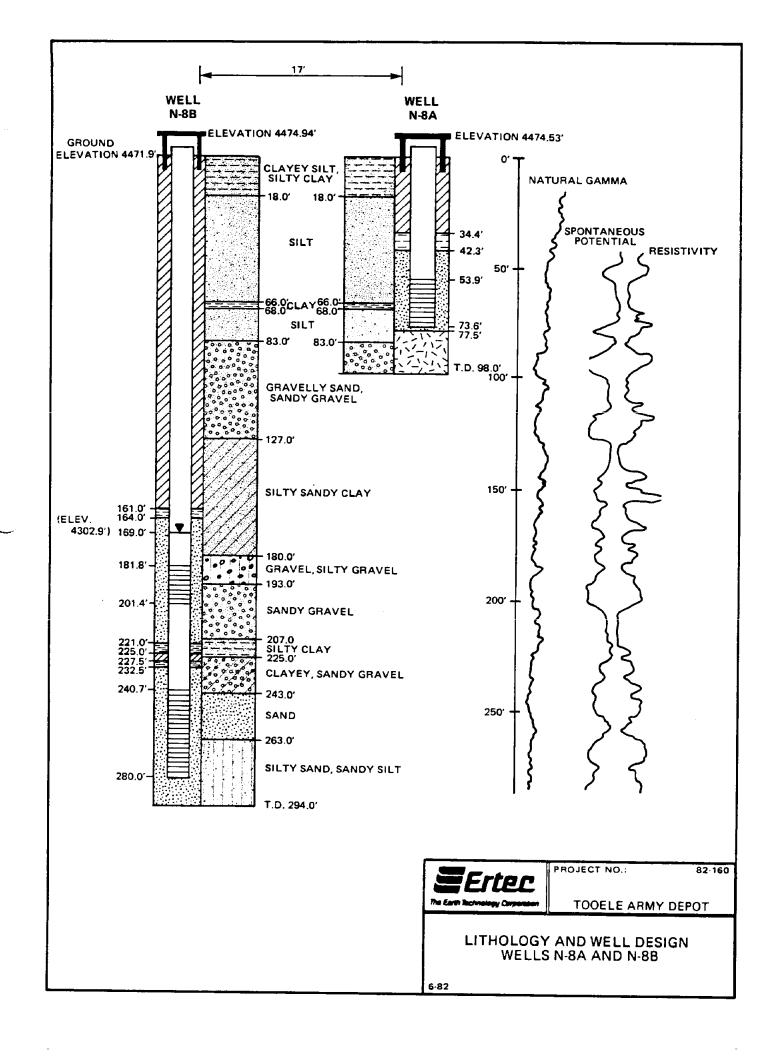


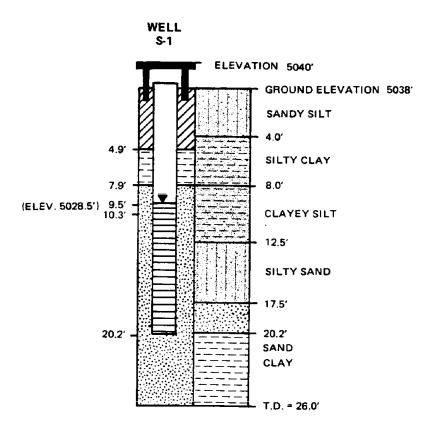


82-160

TOOELE ARMY DEPOT

LITHOLOGY AND WELL DESIGN BORING N-7



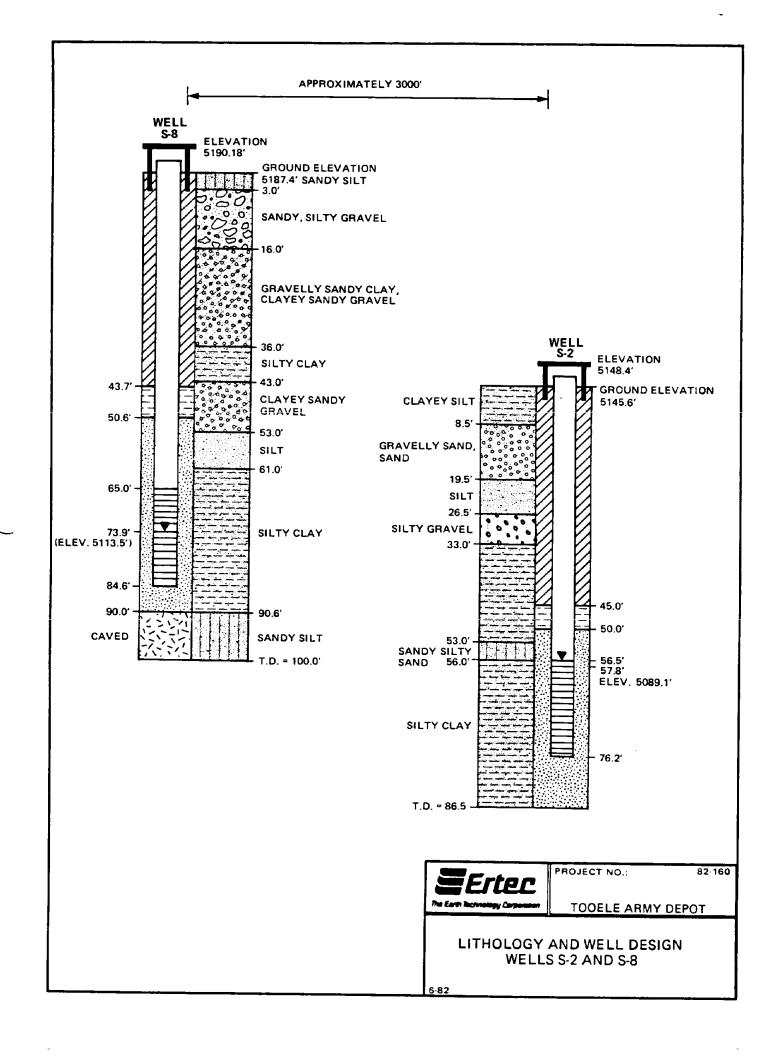


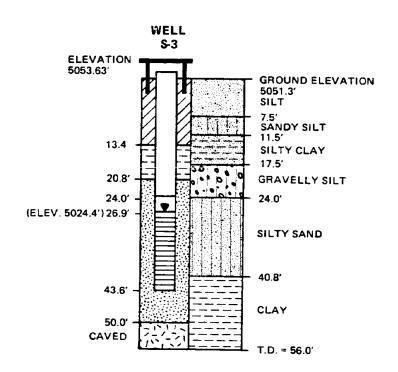


22.16

TOOELE ARMY DEPOT

LITHOLOGY AND WELL DESIGN WELL S-1



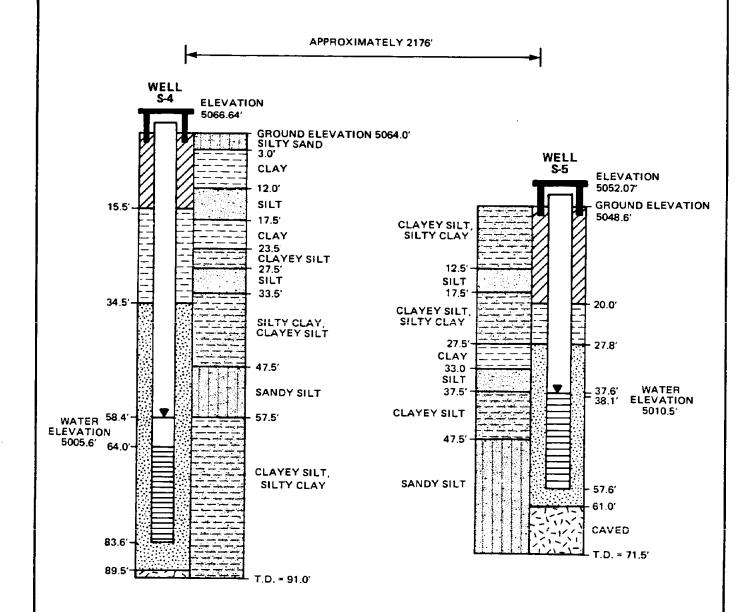




82-160

TOOELE ARMY DEPOT

LITHOLOGY AND WELL DESIGN WELL S-3



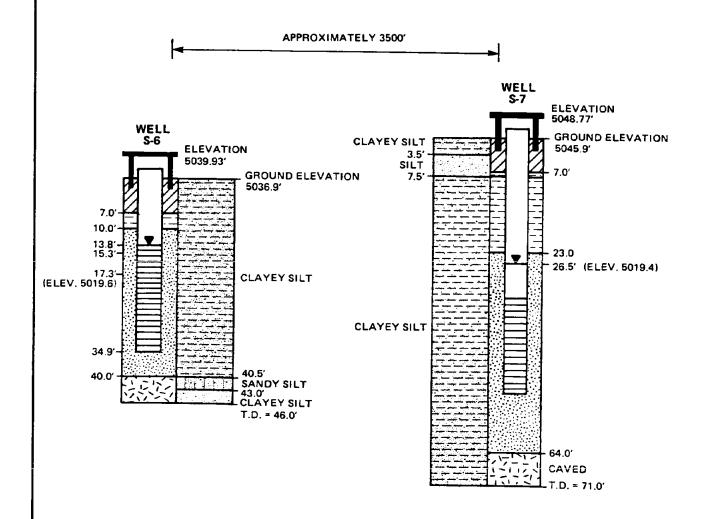


82-160

TOOELE ARMY DEPOT

LITHOLOGY AND WELL DESIGN WELLS S-4 AND S-5

<u>6-82</u>

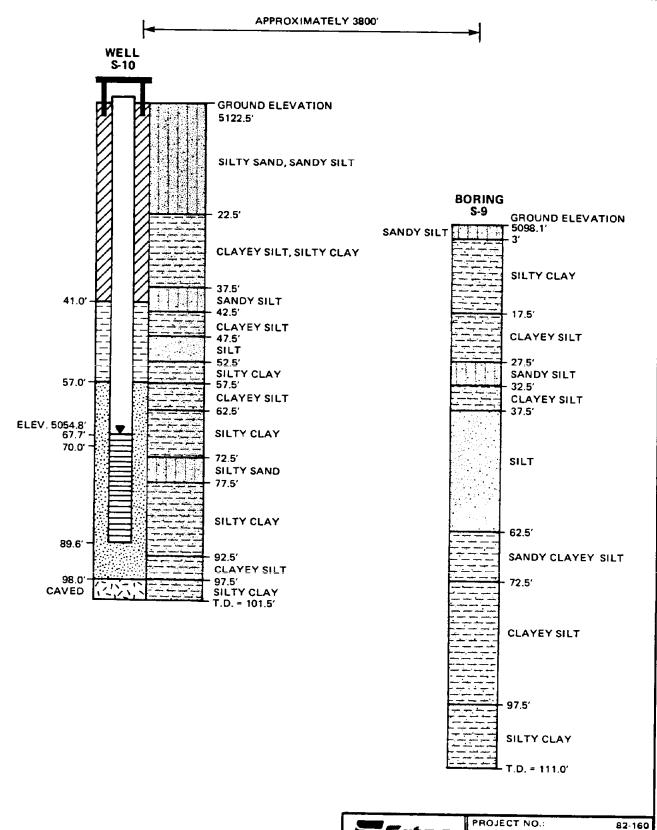




82-160

TOOELE ARMY DEPOT

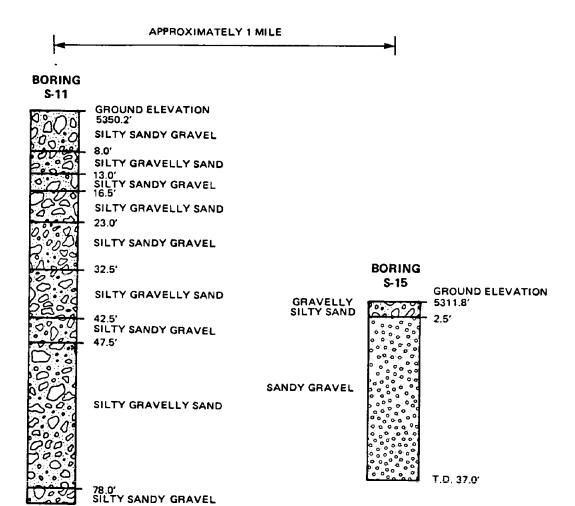
LITHOLOGY AND WELL DESIGN WELLS S-6 AND S-7





TOOELE ARMY DEPOT

LITHOLOGY AND WELL DESIGN WELL S-10 AND BORING S-9



T.D. = 81.0'

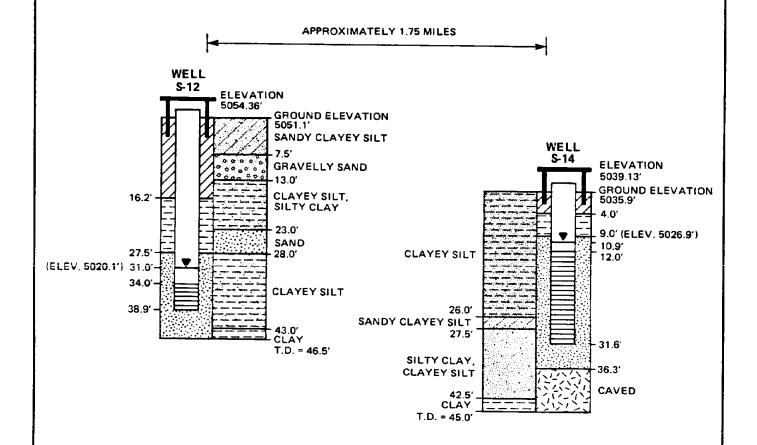


PROJECT NO.:

82.16

TOOELE ARMY DEPOT

LITHOLOGY AND WELL DESIGN BORINGS S-11 AND S-15





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TOOELE ARMY DEPOT

LITHOLOGY AND WELL DESIGN WELLS S-12 AND S-14 Appendix D
Geophysics Program

D-1 Magnetic Surveys

D-1.1 Introduction

Ground magnetic surveys were performed at three potential drilling sites in the North Area of TEAD and four sites in the South Area for the purpose of detecting buried metallic objects, which might pose a hazard to personnel involved in drilling.

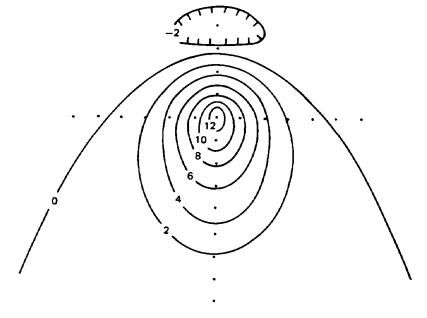
The surveys were designed to detect unexploded ordnance or objects equivalent to a 55-gallon steel drum buried as deeply as 15 feet. The theoretical magnetic anomaly produced by such an object is shown in Figure 1 - a computer simulation for depths of burial of 10 and 15 feet. The simulation assumes that the anomaly is produced solely by magnetization induced by the geomagnetic field - permanent magnetization is not considered. The amplitude of the anomaly is proportional to the mass of the object. For example, a ton of iron buried 15 feet beneath the sensor would create an anomaly of about 500 gammas. Corrosion or rusting may significantly reduce the intensity of an anomaly by reducing the mass of magnetic material.

D-1.2 Survey

The survey areas are squares 80 feet on a side. Measurements were made on a 5-foot grid, producing a total of 289 grid points for each site. The grid was laid out by means of compass, transit, and steel tape. The magnetometer was a Geometrics Model G816 proton precession instrument, which has a resolution of 1 gamma.

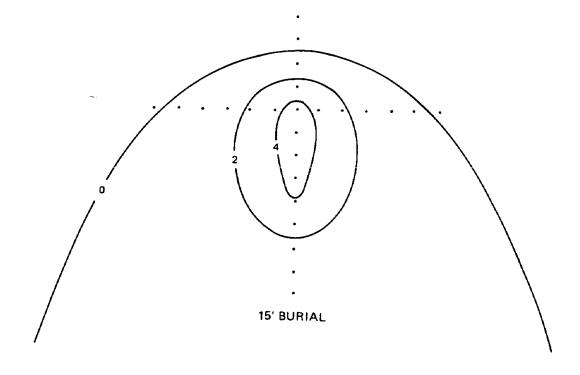
Most magnetometer surveys started in the southwest corner and proceeded northward, making a minimum of two measurements at each grid point. Subsequent traverses were made in alternating southward and northward directions, as

Approved by



10' BURIAL

MAGNETIC INTENSITY UNIT: GAMMAS



EXPLANATION

COMPUTER SIMULATION OF MAGNETIC ANOMALIES FOR A BURIED STEEL DRUM AT TWO DEPTHS OF BURIAL. MAGNETIC SENSOR IS 3' ABOVE GROUND SURFACE. DOTS IN GRID ARE AT 5 SPACING.



PROJECT NO.:

82-160

USATHAMA DRILL SITE SURVEYS

THEORETICAL MAGNETIC ANOMALIES

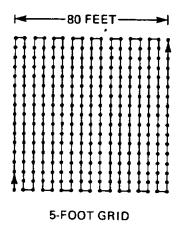
4-82

FIGURE 1

shown in Figure 2. The magnetic sensor was carried in a back pack, with the instrument console supported on the operator's chest by shoulder straps. All magnetic materials were removed from the operator's body, and each measurement was taken while facing in the direction which produced maximum signal strength for the sensor, the response of which is somewhat directional. At each grid point the location, the time, two magnetic readings, and possible comments were recorded. Comments noted the presence of metallic objects, electrical wires, or other relevant facts.

D-1.3 Drift

The geomagnetic field is not constant. It undergoes a diurnal variation of about 50 gammas and has unpredictable variations (micropulsations) of a few minutes duration and a few gammas of amplitude. Magnetic storms are always possible. They can last a day or more and can create amplitude fluctuations of 100 gammas or more. In order to remove such temporal fluctuations (drift) from the survey data, it is necessary to record magnetic intensity at a sationary base station during the survey and subtract the base station values from the survey values. A continuous-recording base station magnetometer, a Geometrics Model G826 proton precession instrument, was operated throughout each survey day. In addition, for surveys S-6, S-7, and S-9, repeated measurements were made at the center of the survey area after each traverse. It was found that the repeated measurements at the center of the survey site gave a significantly better estimate of the drift than the continuous recorder situated a few miles away. The effect of incomplete drift correction shows on some of the low-relief magnetic contour maps as small-amplitude lineations parallel to the survey lines. They do not affect detection capability.





PROJECT NO.:

82-160

USATHAMA DRILL SITE SURVEYS

SURVEY TRAVERSE PATTERN USED FOR ALL SITES

2-82

FIGURE 2

D-1.4 Data Processing

As each site survey was completed, the data were sent to Ertec's office in Long Beach, California, for processing and interpretation. Data processing consisted of coding the data for Ertec's Harris 800 computing system, averaging the magnetometer readings for each grid point, interpolating between drift measurements, subtracting the drift from the survey data, and machine-contouring the processed data. Figures 3 through 9 show the resultant magnetic intensity patterns.

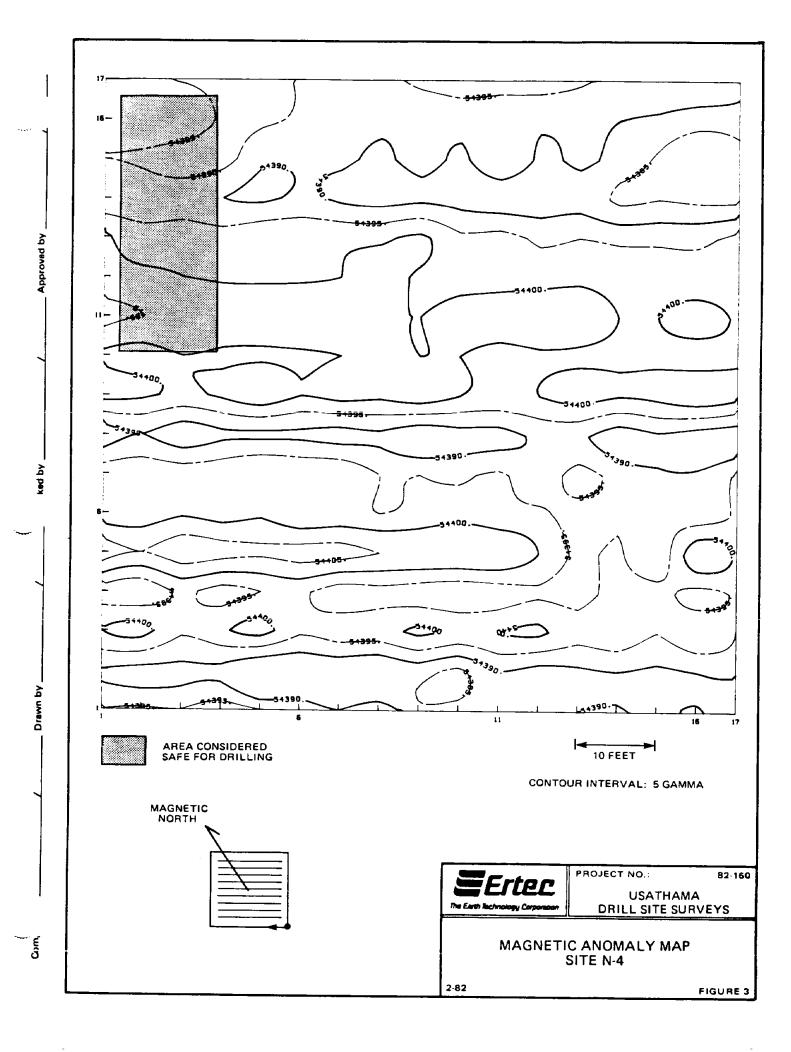
A contour interval of 5 gammas was chosen for all survey sites except N-6NEW, for which a 100 gamma contour interval was necessary to depict the exceptionally large anomalies there. Some sharp corners appear in the contours of the low-relief maps; they are the natural consequence of contouring integer data having only a few units of variation.

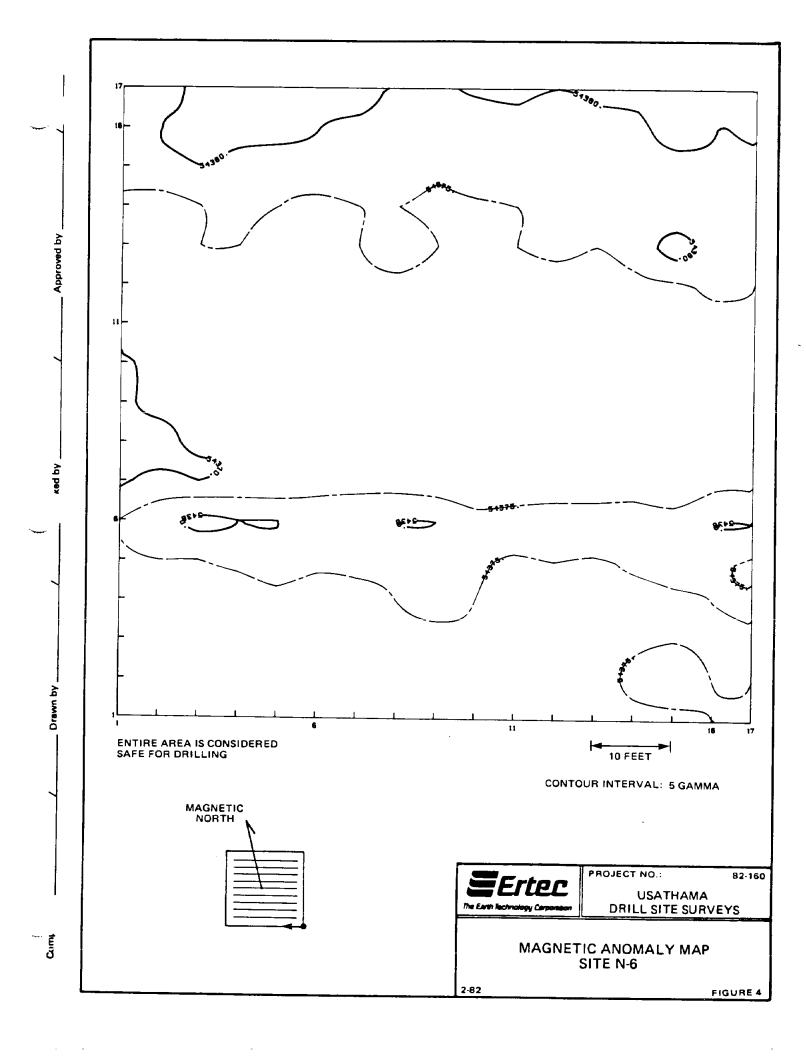
D-1.5 Analysis

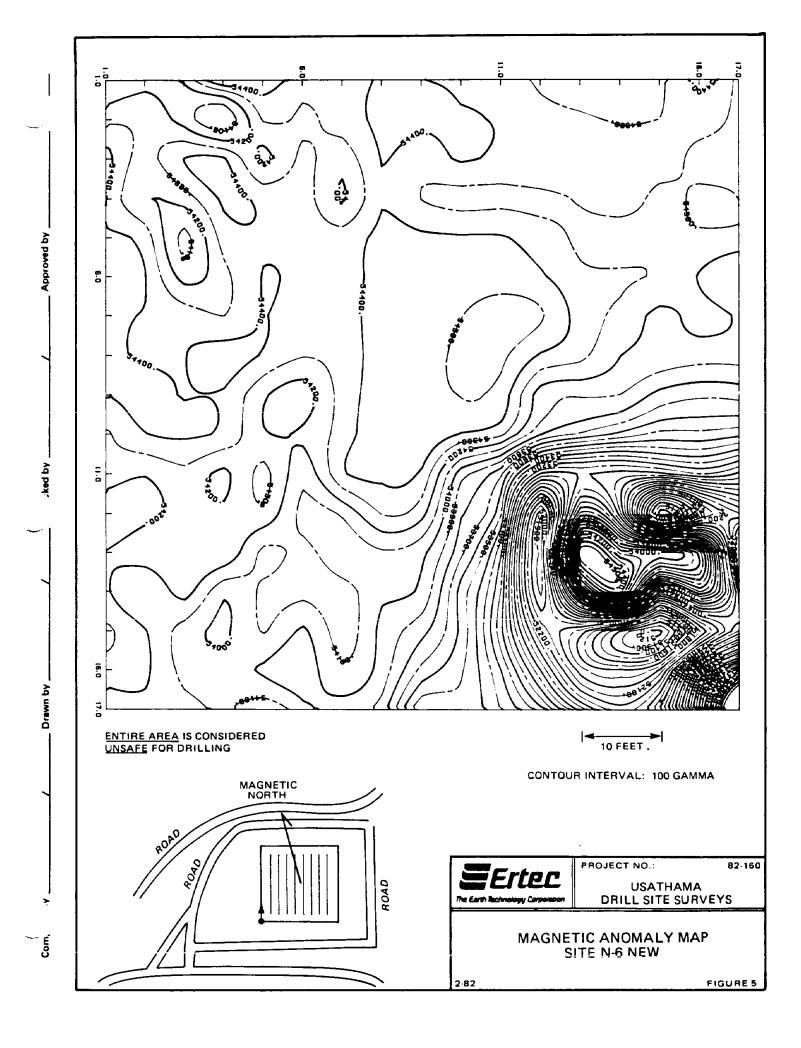
Analysis of the magnetic contour maps for the purpose of selecting a safe drilling site consists simply of avoiding areas of magnetic relief and seeking the flattest and most uneventful areas. The uneventful areas are shown as shaded areas on the maps in Figures 3 through 9.

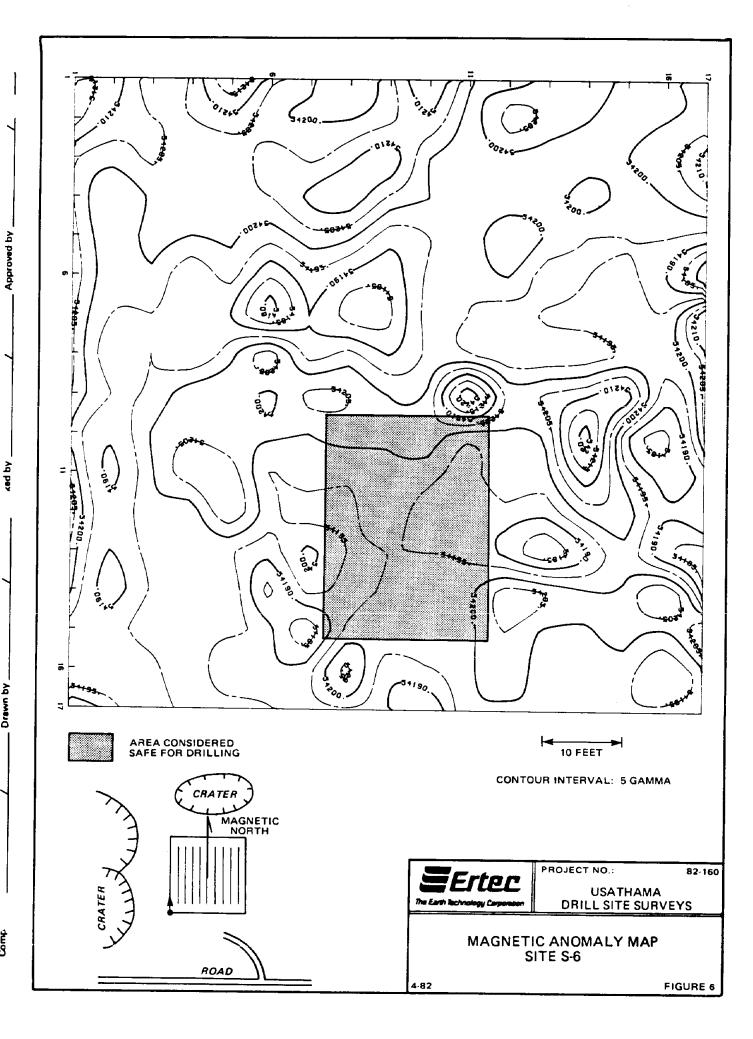
All sites, with the exception of N-6NEW, have drillable areas. Site N-6NEW shows high magnetic relief over the entire area, especially in the southeast corner, where a 3000-gamma anomaly indicated the presence of several tons of iron, or its equivalent.

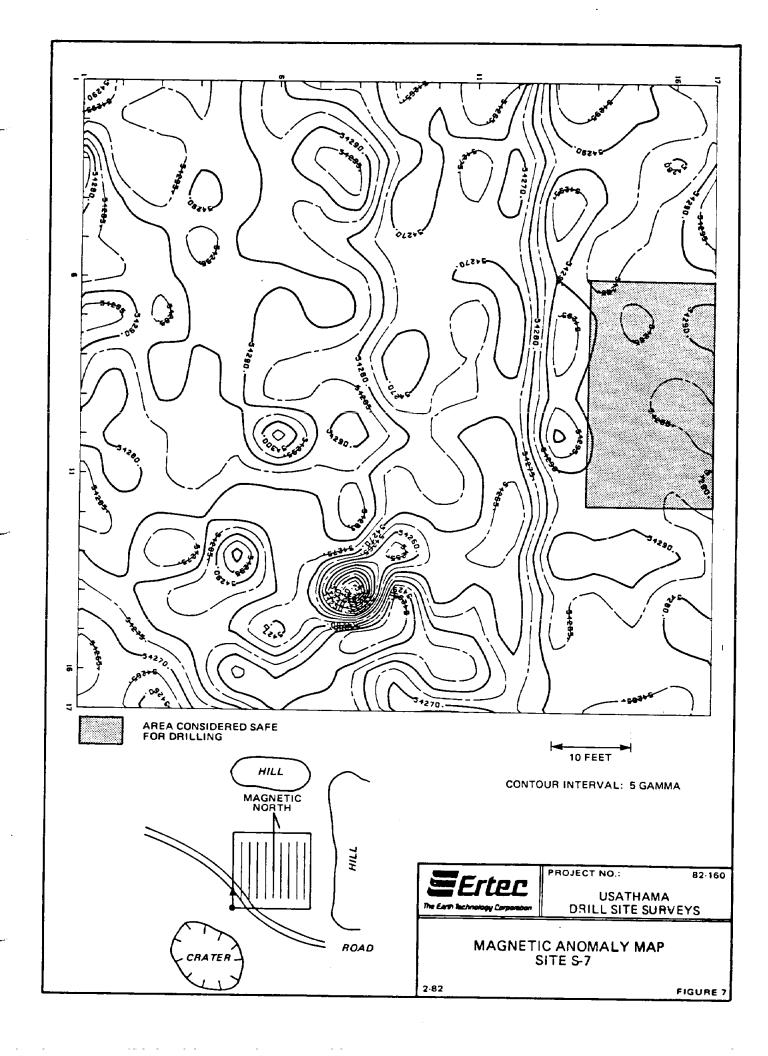
It must be noted that the original site for N-6 was considered inadequate for drilling for reasons unrelated to the magnetometer survey results. After

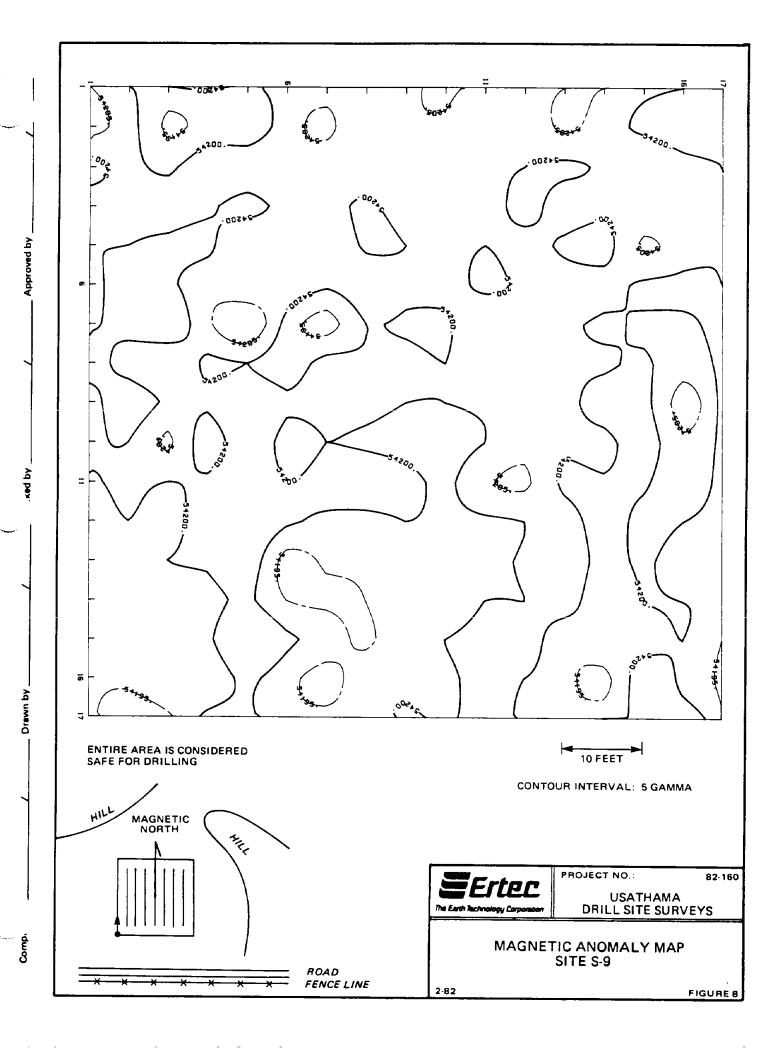


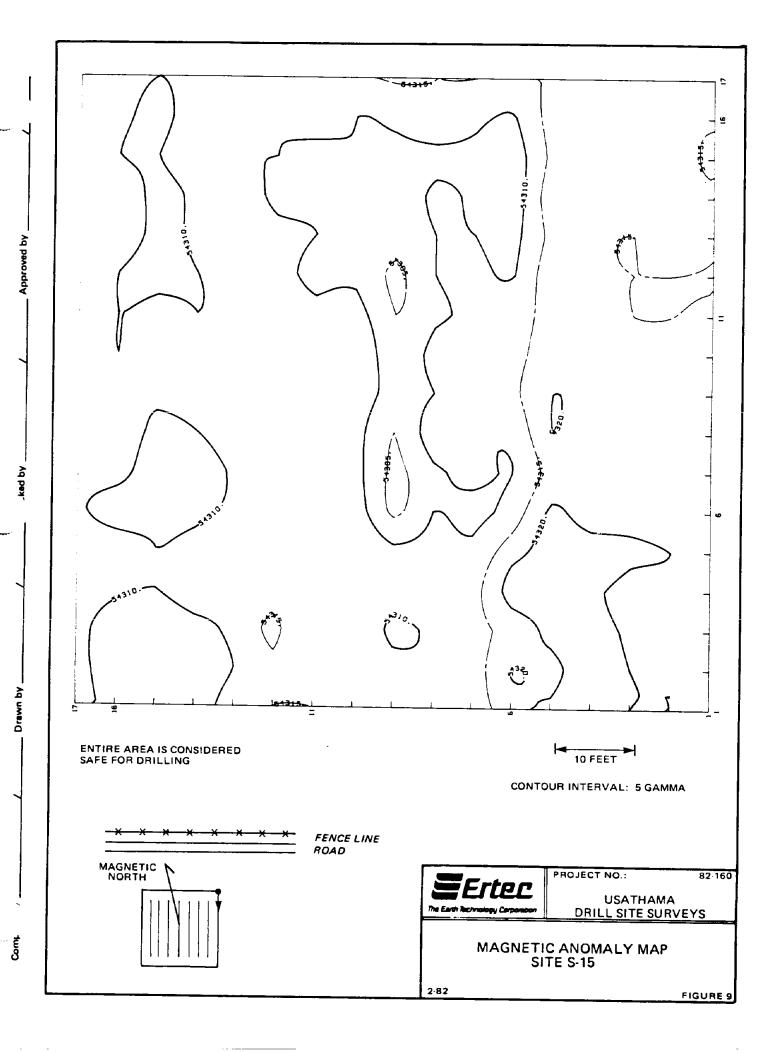












further field investigation upon obtaining new aerial photos of the area, it was decided by Ertec that well N-6 should be re-located further west. Hence, the magnetometer survey for N-6NEW. Since no area could be established that would be safe for drilling, N-6 was again re-located to a more suitable area where little danger of UXO's existed.

D-2 Gravity Survey

D-2.1 Introduction

A gravity survey was conducted at the Tooele Army Depot from the period December 7th through 17th, 1981. The purpose of the survey was to obtain gravity data to provide a conceptual model of the contact between unconsolidated basin fill materials and the underlying bedrock. This model, together with the results of hydrologic studies, would provide information to understand the ground-water flow regime and pathways that transport and distribute potential pollutants. The preliminary gravity results were used to select locations for seismic refraction survey lines and electrical resistivity soundings. Data from the refraction lines corroborated the gravity interpretation.

The gravity survey consists of measurements made at stations. Each field gravimeter measurement was corrected for instrument and tidal drift, as well as latitude, elevation, and terrain effects. Gravimeter units were related to absolute gravity by repeated measurements at Clover Base in nearby Clover, Utah (Cook, et. al., 1971). The gravity at Clover Base is given as 979,698.71 milligals and we have calculated that the Simple Bouguer Anomaly there is -202.49 milligals.

The gravity station locations are given as coordinates on the TEAD survey grid (a network of pre-1942 monuments), for example: x = -14,300 ft, y = 15,600 ft, locates a station west and south of the coordinate origin at the east-central boundary of the Depot. Elevations and locations, are inscribed on the survey monuments.

The data were collected by a two-man field crew with a pickup truck, gravimeter, and transit. Most gravity measurements were taken at survey monuments.

For those stations, surveying was unnecessary. There was some difficulty in
searching out monuments hidden in tall grass, and some monuments were missing.

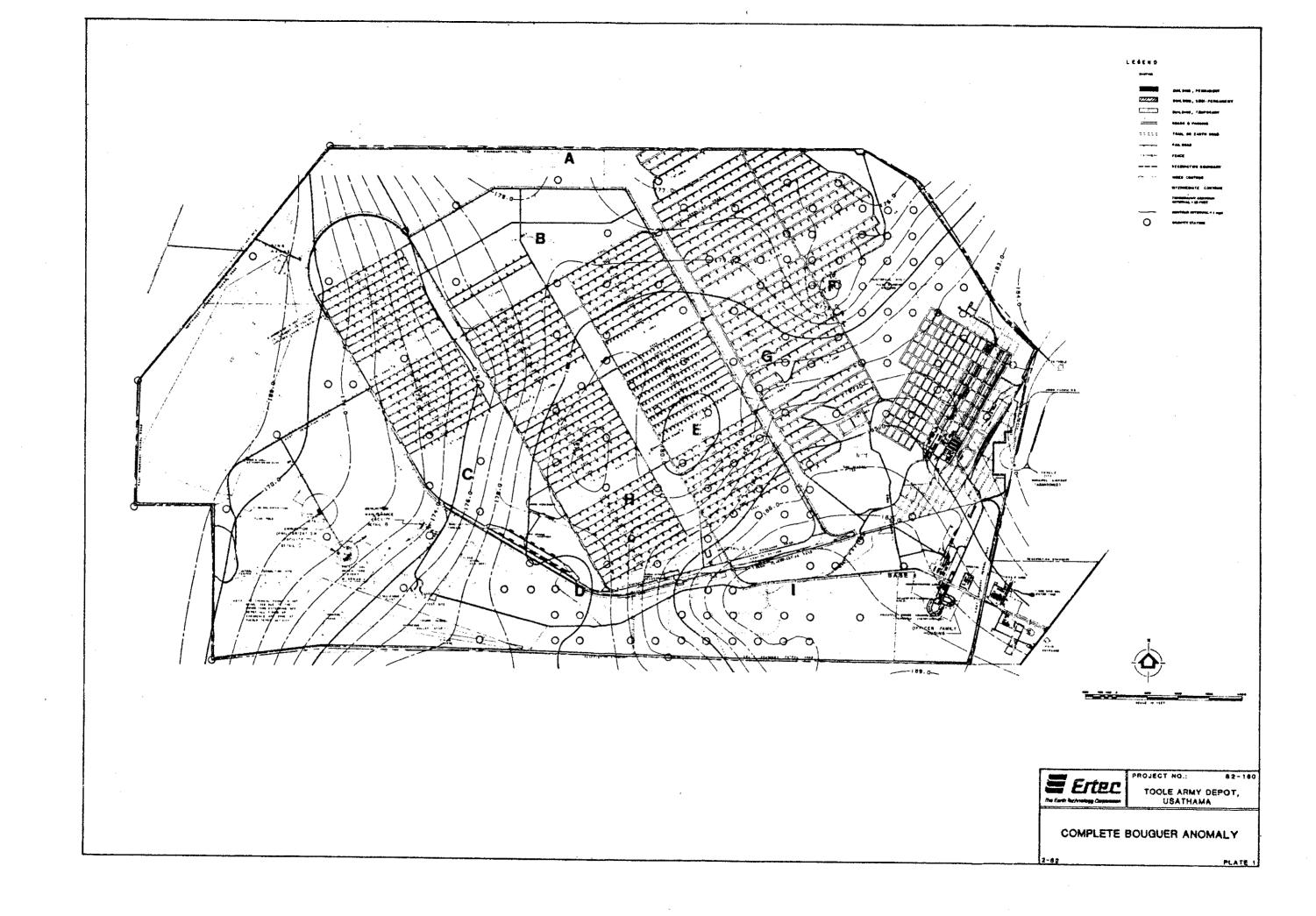
Where there were no monuments, measurements were made at points that could be located on the base map, or located from such points by transit survey.

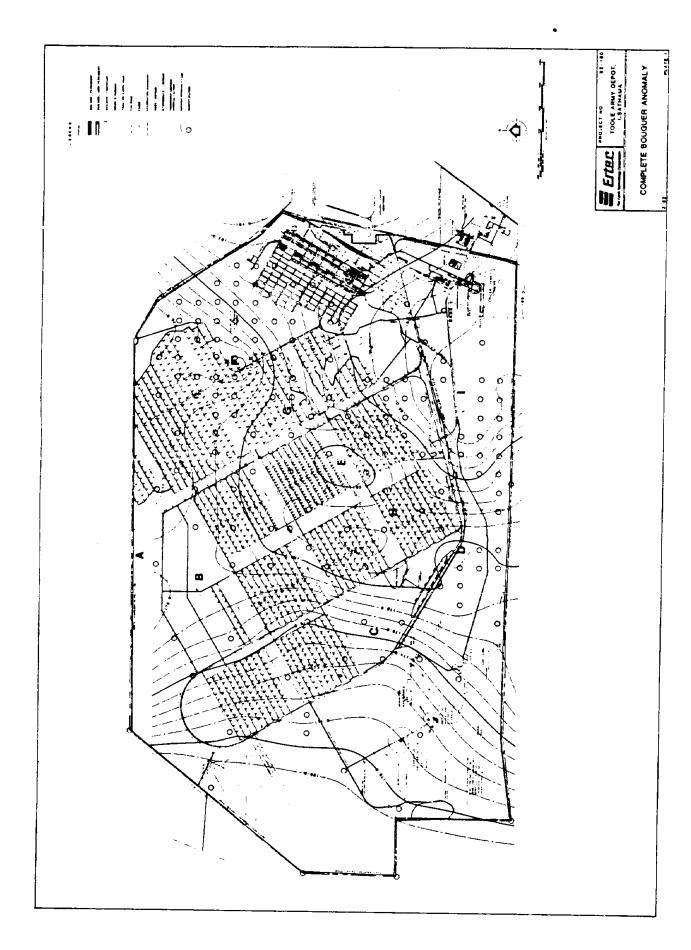
Elevations for these points were taken from the contours of a topographic map.

D-2.2 Analysis of the Tooele Gravity Data

A contour map of terrain-corrected Bouguer (gravitational) Anomaly (Complete Bouguer Anomaly or CBA) is shown in Plate 1. Terrain corrections were done for a distance of 32,490 feet (the equivalent of Zone K on the Hammer Chart; see for example: Dobrin, 1976). Corrections for greater distances from the station were not done because the correction factor became the same for every station within the survey area. The terrain-corrected data were gridded using a minimum curvature algorithm (Briggs, 1974; Swain, 1976) and contoured by computer. This algorithm computes the surface of least curvature which passes through an aribtrarily distributed set of data points and produces an interpolated set of values on a uniform grid suitable for machine contouring. The grid spacing was chosen to be slightly less than the average data spacing.



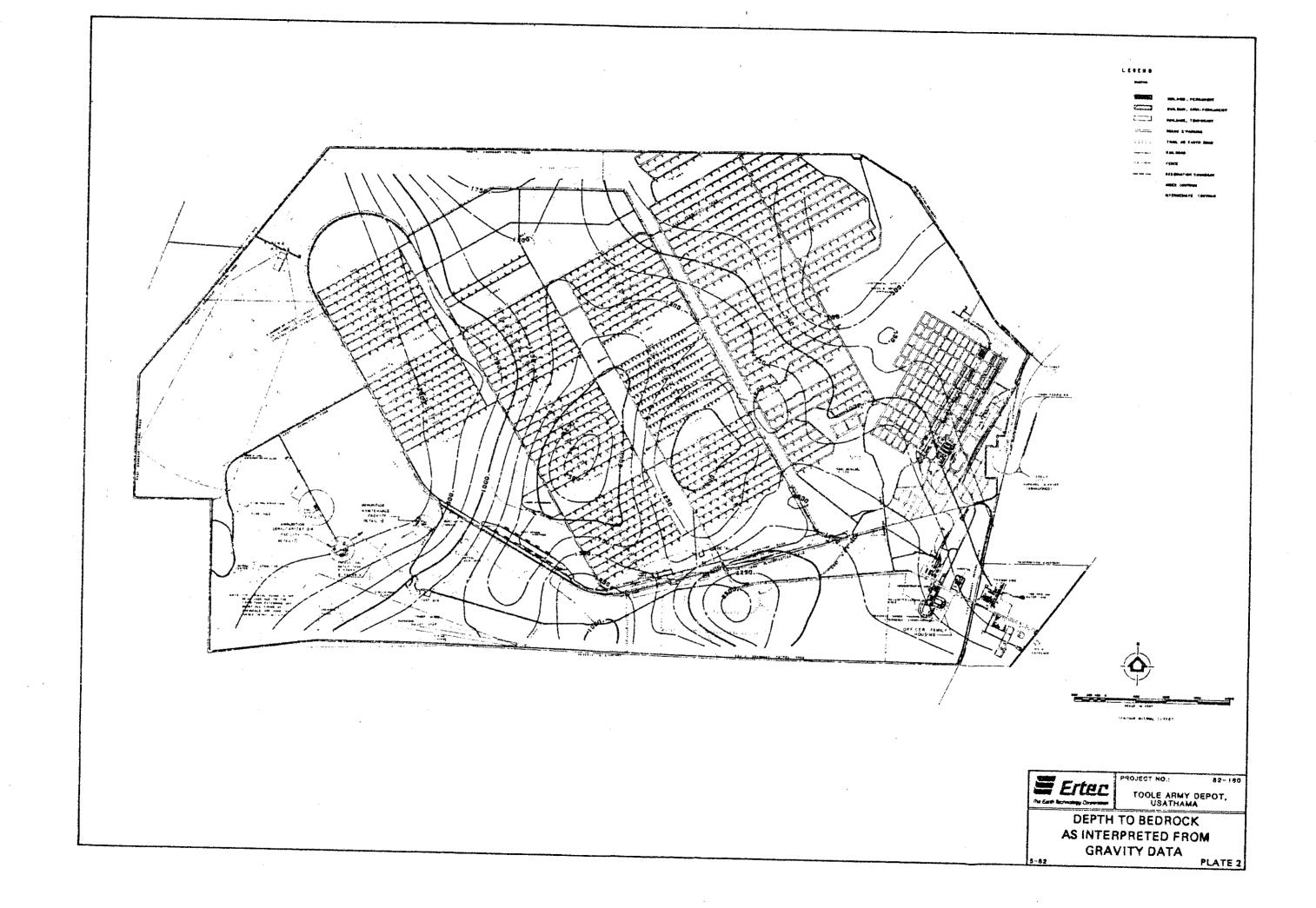




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four separate reductions, to account for four geometrical effects, are made to the observed gravity at each station to arrive at its Bouguer Anomaly value.

a. Free-Air Effect: Gravitational attraction varies inversely as the square of the distance from the center of the earth. Thus, corrections must be applied for elevation. Observed gravity levels are corrected for elevation using the normal vertical gradient of:

FA = 0.09406 mg/ft (-0.3086 milligals/meter)

where FA is the free-air effect (the rate of change of gravity with distance from the center of the earth). The free-air correction is positive in sign since the correction is opposite the effect.

b. Bouguer Effect: Like the free-air effect, the Bouguer effect is a function of the elevation of the station, but it considers the influence of a slab of earth materials between the observation point on the surface of the earth and the corresponding point on the geoid (sea level). Normal practice, which is to assume that the density of the slab is 2.67 grams per cubic centimeter, was followed in these studies. The Bouguer correction (B_C), which is opposite in sign to the free-air correction, was defined according to the following formula.

 $B_C = 0.01276 (2.67) \text{ hf (milligals per foot)}$

 $B_C = 0.04185 (2.67) h_m \text{ (milligals per meter)}$

where h_{f} is the height above sea level in feet and hm is the height in meters.



- c. Latitude Effect: Points at different latitudes will have different values of gravity to two reasons. The earth (and the geoid) is spheroidal, or flattened at the poles. Since points at higher latitudes are closer to the center of the earth than points near the equator, gravity at the higher latitudes is larger. As the earth spins, the centrifugal accerlation causes a slight decrease in the measured value of gravity. At the higher latitudes where the earth's circles of latitude are smaller, the centrifugal acceleration diminishes. The gravity formula for the Geodetic Reference System, 1967, gives the theoretical value of gravity at the geoid as a function of latitude. It is:
 - $g = 978.0281 (1 + 0.0053204 \sin^2 0 0.0000058^2 20) gals$

where g is the theoretical acceleration of gravity and o is the latitude in degrees. The positive term accounts for the spheroidal shape of the earth. The negative term adjusts for the centrifugal acceleration.

The previous two corrections (free air and Bouguer) adjust the observed gravity to the value it would have at the geoid (sea level). The theoretical value at the geoid for the latitude of the station is subtracted from the adjusted observed gravity and the remainder is called the Simple Bouguer Anomaly (SBA). Most of this represents the effect of material beneath the station, but part of it may be due to irregularities in terrain (upper part of the Bouguer slab) around the station.

d. Terrain Effect: Topographic relief around the station has a negative effect on the gravitational force at the station. A nearby hill has

upward gravitational pull and nearby valley contributes less downward attraction than a nearby material would have. Therefore, the corrections are always positive. Corrections are made to the SBA when the terrain effects are 0.1 milligal or larger. Terrain corrected Bouguer values are called the Complete Bouguer Anomaly (CBA). When the CBA is obtained, the reduction of gravity at individual measurement points (stations) is complete.



D-2.5 Method of Interpretation

D-2.5.1 Regional - Residual Separation

To interpret the gravity data, the portion of the CBA that might be caused by the light-weight, basin-fill material must be separated from that caused by the heavier bedrock material which forms the surrounding mountains and presumably the basin floor. The first step is to estimate a regional field. This is an estimation of the values the CBA would have if the light-weight sediments (the anomaly) were not there. Since the valley-fill sediments are absent at the stations read in the mountains, one approach is to use the CBA values at bedrock stations as the basis for constructing a second-order polynomial surface to represent a retional field over the valley.

Where there are insufficient bedrock stations to define a satisfactory regional trend, another approach is to estimate the regional by the process of upward continuation of the CBA field. A principal result of potential field theory is that a field quantity satisfying Laplace's equation in a three-dimensional volume of space is specified completely by the value it has on the surface bounding that volume (Grant and West, 1965). Since the gravitational field statisfies Laplace's equation, its value anywhere above the surface of the earth can be found using only the value of gravity on the surface of the earth, regardless of the mass distribution that produces the value of gravity in the first place. On this basis, the Bouguer anomaly is readily continued to level surfaces above the ground.

An important property of upward continuation is that the resultant field (which can be represented by a contour map), changes more with respect to

shallow sources than it does with respect to deeper souces. The anomalous parts of the field ascribed to shallow density distribution tend to vanish as the continuation is carried upward; whereas the field produced by deeper sources changes only slightly, so that upward continuation produces regional-type fields.

The difference between the CBA and the regional field is called the residual field or residual anomaly. The residual field is the interpreter's estimation of the gravitational effect of the geologic anomaly.

D-2.5.2 Interpretation of the Residual Anomaly

If the regional is well chosen, the magnitude of the residual anomaly is a function of the thickness of the anomalous (fill) material and the density contrast. The density contrast is the difference in density between the alluvial and bedrock material. If this contrast were known exactly, an accurate calculation of the thickness could be made. Generally, the densities are not well known and vary within the study area. Therefore, it is necessary to use densities typical of materials similar to those in the study area.

If the selected average density contrast is smaller than the actual density contrast, the computed depth to bedrock will be greater than the actual depth and vice-versa. The computer depth is inversely proportional to the density contrast. A ten percent error in density contrast produces a ten percent error is computed depth.

Once the density contrast betwen the alluvium and bedrock is established, there are several methods available for determining the form of the alluvium-

bedrock interface. One way is to use an interative computer porogram which will yield some simple model of the interface approximately explaining the residual gravity anomaly (Dordell, 1970). An alternative approach is to assume the form of the interface a priori and calcuate what effect this would have on the residual gravity anomaly. By continually adjusting the model, one may obtain a reasonable estimate of the interface. There are computer programs that will calculate the gravitational effect of two-dimensional (Talwani, et al., 1959) and three-dimensional (Cordell, 1970; Plouff, 1975) bodies.

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Gravity Survey Data

SIMPLE AND COMPLETE BOUGUER ANOMALIES ASSUMING A DENSITY OF 2.67 GRAVITY SURVEY DONE AT TOOELE, UTAH

X, Y= POSITIONS ON LOCAL COORDINATE SYSTEM (FEET) T=TERRAIN CORRECTION

				1 1 - 1111111	N CORRECTION
ID	x	Y	SBA	T	CBA
1	-6870, 000	-10670, 000	-192. 580	יירי כ	100 054
2	-16900, 000	-11700.000	-191. 013	3. 327	-189. 254
3	-16900.000	-13000.000	-191. 001	2. 600	-188. 413
4	-15600.000	-13000.000	-192. 424	2. 541	-188. 460
5	-14300.000	-13000.000	-193. 245	2.513	-189. 911
6	-13000.000	-13000.000	-173. 109	2. 558	-190. 687
フ	-18200.000	-13000.000	-189. 394	2. 649 2. 544	-190. 460
8	-19500, 000	-14300.000	-188. 245	2. 566 3. 813	-186. 828
9	-18200.000	-14300.000	-188. 615	2. 812 2. 932	-185. 433
10	-16900.000	-14300.000	-189. 831		-185. 684
11	-15600,000	-14300.000	-191, 439	3.004	-186. 827
12	-14300.000	-14300.000	-192. 498	2. 845 2. 853	-188. 594
13	-13000.000	-14300.000	-192. 729	2.852	-189. 646
14	-11700.000	-14300.000	-192. 788	2.899	-189. 830
15	-26000,000	-10400.000	-184. 063	3.088	-189. 700
16	-24700.000	-11700.000	-182. 937	2. 560	-181. 503
17	-23400,000	-13000.000	-184. 263	2. 608	-180. 329
18	-23400.000	-14300,000	-185. 044	2.826	-181. 438
19	-20800, 000	-14300.000	-186. 425	2. 937	-182.107
20	-9100.000	0.000	-184. 871	2. 962	-183. 463
21	-9100.000	2600.000	-181. 638	1.869	-183.002
22	-9100.000	3900.000	-179. 893	1.538	-180. 100
23	-7800, 000	0. 000	-185. 254	1.395	-178. 498
24	-10400, 000	2600, 000	-181, 092	2. 058	-183. 196
25	-10400.000	3900.000	-181. 072 -179. 024	1. 681	-179. 411
26	-11700.000	5200. 000	-177. 024 -179. 187	1. 543	-177. 481
27	-9100.000	6500. 000	-177. 187 -179. 187	1.658	-177. 529
28	-9100.000	7800, 000	-177. 187 -178. 304	1.454	-177, 733
29	-7800, 000	6500. 00C	-178. 304 -179. 282	1. 419	-176. 885
30	-6500, 000	6500.000	-177. 262 -180. 614	1.521	-177. 761
31	-9100.000	5200. 000	-178. 891	1.327	-179, 287
32	-10400.000	3900.000	-176. 839	1.419	-177. 473
33	-13000.000	5200.000	-176. 639 -179. 872	1. 543	-175. 296
34	-14300.000	3900.000	-177. 672 -181. 320	1. 465	-178. 407
35	-7800, 000	2600.000	-183. 410	1.521	-179. 799
36	-6500.000	2600, 000	-183, 410 -183, 864	1.579	-181.831
37	-3900, 000	2600.000	-185. 382	1.607	-182. 257
38	-7800. 000	1300.000	-184. 088	1. 676	-183, 706
39	-7800.000	5200.000		1.772	-182. 316
40	-7800, 000	7800. 000	-180. 191 -178. 180	1.416	-178.775
4 i	-6500.000	7800, 000	-178. 190 -179. 542	1. 497	-176. 693
42	-6500.000	5200.000	-179, 542 -181, 736	1. 274	-178. 268
43	-10400.000	9100.000	-181, 736 -178, 015	1.376	-180.360
44	-11700.000	7800.000	-178. 853	1.489	-176. 526
45	-13000.000	6500, 000	-179. 637	1. 581	-177. 272
46	-11700, 000	6500.000	-179. 214	1. 457	-178. 180
47	-10400.000	7800, 000	-179. 214 -178. 685	1.582	-177. 632
	· 	. 255, 500	170.003	1. 457	-177. 228

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48	-16900, 000	5200. 000	-181. 172	1. 780	-179, 392
49	-13000.000	-2600.000	-186. 338	2. 074	-184. 264
50	-13000.000	0.000	-183. 546	2. 051	-181. 495
51	-15600.000	0.000	-182. 965	1. 770	-181.195
52	-15600, 000	2600.000	-182. 597	1. 690	-180. 907
53	-14300.000	-1300.000	-184. 209	1. 993	-182.216
54	-13000.000	1300.000	-182. 761	1. 724	-181.037
55	-11700.000	3900.000	-179. 654	1. 683	-177. 971
56	-13000,000	3900.000	-180. 617	1. 552	-179.065
57	-19500.000	6500.000	-179. 822	1. 422	-178. 400
58	-15600,000	7800.000	-179. 527	1. 374	-178. 153
59	-13000, 000	9100.000	-179. 278	1. 375	-177. 903
60	-19500.000	9100.000	-17B. 197	1. 344	-176.853
61	-18200.000	7800, 000	-179. 039	1. 325	-177. 714
62	-15600.000	-11700.000	-192.312	2. 520	-189. 792
63	-10400.000	-10400.000	-192. 623	2. 652	-189.971
64	-9 100.000	-9 100.000	-191.722	2. 677	-189. 045
65	-11700.000	-6500, 000	~189.061	2. 495	-186. 566
66	-11700.000	-5200.000	-189. 515	2. 465	-187. 050
67	-10400, 000	-2600, 000	-187. 871	3. 350	-184. 521
68	-7800, 000	-2600, 000	-188. 245	2. 219	-186. 026
69	-5200, 000	2600, 000	-184. 467	1.621	-182. 846
70	-22100, 000	6500, 000	-180. 113	1. 386	-178. 727
71	-24700, 000	3900.000	-181.110	1. 593	-179. 517
72	-22100, 000	3900.000	-181. 087	1. 523	-179, 564
73	-24700, 000	1300.000	-181. 624	1. 699	-179, 925
74	-23400, 000	-1300.000	-183. 146	1. 978	-181, 168
75	-22100.000	0. 000	-183. 268	1.686	-181.582
76	-22100.000	1300.000	-182. 533	1. 630	-180, 903
77	-19500.000	3900. 000	-181. 764	1.541	-180, 223
78	-18200.000	2600.000	-182. 538	1.569	-180. 969
79	-18200.000	0. 000	-182. 937	1. 765	-181. 172
80	-20800.000	-1300. 000	-183. 912	1.841	-182.071
81	-16900.000	-2600.000	-182. 412	2. 026	-180. 386
82	-19500.000	~ 6500. 000	- 184. 163	2. 371	-181. 792
83	-20800.000	- 9100.000	-185. 733	2. 517	-183. 216
84	-15600.000	- 7800. 000	-187. 309	2. 339	-184, 970
85	-14300.000	-6 5 00. 000	-187. 784	2. 459	-185. 325
86	-16900.000	~6500.000	-184. 891	2. 611	-182. 280
87	-15600,000	-5200.000	-184. 915	2. 240	-1B2. 675
88 89	-18200.000	-5200.000	-182. 942	2. 554	-180. 388
90	-14300, 000	-3900.000	-185. 711	3. 301	-182. 410
70 91	-19500.000 -77100.000	-2600.000	-183. 563	2. 436	-181. 127
92	-22100, 000 -24700, 000	-6500.000	-185. 280	2. 339	-182. 941
93	-19500, 000 -19500, 000	-5200.000	-184. 476	2. 163	-182. 313
73 94	-14500, 000 -16900, 000	-10400.000	-187. 076	2. 563	-184. 513
7 4 95	-13000.000 -13000.000	-9100.000	-187, 937	3.009	-184. 928
96	-5200.000 -5200.000	6500, 000	-189, 410	2. 414	-186, 996
97	-6500.000 -6500.000	3700.000	-184. 338	1. 513	-182. 825
98	-3900. 000 -3900. 000	3900. 000 3900. 000	-183. 150	1.466	-181. 684
99	-5200. 000 -5200. 000	5200. 000 5200. 000	-185. 141	1. 575	-183. 566
100	-7800. 000	3900.000	-183. 105 -181. 701	1. 415	-181. 690
101	-14300.000	-7800. 000 -7800. 000	-181. 701 -188. 775	1. 478	-180. 223
102	-13000.000	-7800. 000 -7800. 000	-188. 775 -180. 304	2. 668	-186. 107
102	10000.000	7000. 000	−19 0. 3 06	2. 512	-187. 794

103	-13000. 000	-9100.000	_101 252	2 400	
104	-11700.000	-7800, 000 -7800, 000	-191. 353 -190. 7/3	2. 499	-188. 854
105	-22100.000	-3900.000	-190, 763 -184, 984	2. 594	-188. 169
106	-24700.000	-2600.000	-182. 950	2.002	-182. 983
107	-28600.000	-1300.000	-177. 5 95	1. 945	-1B1. 005
108	-29900.000	3900.000	-177, 575 -178, 800	1 969	-175. 626
109	-36400.000	3900.000	-178. 800 -172. 911	1. 663	-177. 137
110	-32500.000	0.000	-175. 717	1.849	-171.062
111	-31200.000	-3900.000	-176. 724	1. 926	-173. 791
112	-28600, 000	-7800.000	-180. 427	2. 242 2. 296	-174. 482
113	~28600.000	-5200 000	-179. 288	2. 276 2. 225	-178. 131
114	-31200.000	-6500.000	-176. 652	2. 263	-177. 063 -174. 200
115	-22100.000	-10400.000	-184. 134	2. 534	-174. 389 -181. 480
116	-24700, 000	9100.000	-180,006	1. 400	-181.600
117	~29900.000	7800.000	-179. 607	1. 576	-178. 606 -178. 600
118	-9 100. 000	-13000.000	-193. 249	3. 201	-178.032
119	-11700.000	-13000.000	-192. 974	3. 201 2. 773	-190. 04B
120	-11700.000	-10400.000	-192. 492	2. 773 2. 629	-190. 201
121	-6500, 000	-7800.000	-191. 719	2. 874	-189. 863
122	-1340.000	128.000	-186. 904	2. 024	-188. 845
123	-38932.000	-4030, 000	-173. 990	2. 101	-184. 880
124	-41120.000	-8080.000	-175. 556	2. 169	-171.889 -172.007
125	-46540, 000	-7410.000	-171. 810	2. 444	-173. 387 -148. 344
126	-46260, 000	-964.000	-170. 599	2. 423	-169. 366 -179. 137
127	-36608, 000	10776. 000	-175. 250	1.842	-168. 176 -173. 400
128	-40300, 000	5200.000	-170. 545	1. 924	-173. 40B
129	-42511.000	-15334.000	-176. B76	3. 200	~168. 621 ~172. 474
130	-28600, 000	-14300.000	-183. 324	2. 969	-173. 676
131	-11700.000	0.000	-183. 906	1. 937	-180. 355 -181. 848
132	-10400. 000	5200, 000	-178. 545	1. 51G	-181. 969 -177. 035
133	-9 100. 000	5200 . 000	-178. 847	1. 419	-177. 035 -177. 429
134	-13000.000	5200, 000	-179. 834	1. 465	-177. 427 -178. 369
135	-14300.000	5200.000	-180. 600	1. 466	-178. 387 -179. 134
136	-15600, 000	5200. 000	-180. 962	1. 443	-179. 519
137	-10400.000	1300. 000	-183, 118	1.860	-181. 258
138	-14300.000	3900.000	-181. 384	1. 521	-179. B63
135	-5312.000	0.000	-186. 210	1.859	-184. 351
140	-3660. 000	1300.000	-185. 336	1. 778	-183. 558
141	-4604. 000	-1117. 000	-186. 5 89	2. 044	-184. 543
142	-6223. 000	-2454.000	-188. 184	2. 184	-1B6. 000
143	-3084. 000	-2704, 000	-187. 123	2. 772	-184. 351
144	-31200.000	-9 100,000	-178. 139	2. 334	-175. 803
145	-35655. 000	- 9696, 000	-176. 768	2. 198	-174. 570
146	-26000.000	-11700.000	-183. 743	2. 607	-181. 136
147	-27300. 000	-11700.000	-183. 777	2. 592	-181. 185
148	-24700.000	-13000.000	-183. 250	2. 754	-180. 496
149	-24700. 000	-14300.000	-184. 563	2. 830	-181.733
150	-32177.000	-11879.000	-180.778	2. 400	-17B. 37B
151	-32262.000	6336. 000	~177. 013	1.608	-175. 405
152	-34355. 000	-1568. 000	-175. 299	2. 050	-173. 249
153	-18886. 000	-15115 000	7187, 909	2. 841	-185 048

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SIMPLE AND COMPLETE BOUGUER ANOMALIES ASSUMING A DENSITY OF 2.3 SURVEY AT TOOELE UTAH.

X, Y= LOCATIONS ON LOCAL COORDINATE SYSTEM T=TERRAIN CORRECTION

	•			•	
ID	X	YY	SBA	T	CBA
1	- 6870. 000	-10670, 000	-190. 755	3. 327	-187. 429
2	-1 6900, 000	-11700.000	-189. 463	2. 600	-186. 863
3	-16900.000	-13000.000	-189. 154	2. 541	-186. 613
4	-15600.000	-13000.000	-190. 598	2. 513	-188.085
5	-14300.000	-13000.000	-191, 435	2. 558	-188. 877
6	-13000.000	-13000.000	-191. 310	2. 649	-188. 661
7	-18200.000	-13000.000	-187. 524	2. 566	-184, 958
8	-19500.000	-14300.000	-186. 013	2. 812	-183. 201
9	-18200.000	-14300.000	-186, 565	2. 932	-183. 634
10	-16900,000	-14300.000	-187. 927	3. 004	-184. 923
11	-15600.000	-14300.000	-189. 549	2. 845	-186. 704
12	-14300.000	-14300.000	-190. 625	2. 852	-187. 773
13	-13000.000	-14300.000	-190. B61	2.899	-187. 962
14	-11700.000	-14300.000	-190. 912	3. 088	-187, 824
15	-26000.000	-10400.000	-182. 737	2. 560	-180. 177
16	-24 700, 000	-11700.000	-181. 440	2. 608	-178. 832
17	-23400.000	-13000.000	-182. 550	2. 826	-179. 725
18	-23400.000	-14300.000	-182. 976	2. 937	-180. 039
19	-20800.000	-14300.000	-184. 268	2. 962	-181. 306
20	-9100. 000	0.000	-183. 870	1.869	-182.001
21	-9100. 000 -9100. 000	2600. 000	-180. 663	1. 536	-179. 125
55	-9100. 000 -9100. 000	3900.000	-178. 901	1. 395	
23	-7800. 000 -7800. 000	0.000	-184. 102		-177. 506 -183. 044
24	-10400.000	2600, 000	-184. 10 <u>2</u> -180. 289	2. 058	-182.044
25 25	-10400.000	3900.000	-160. 264 -178. 264	1. 681	-178. 608
26 23	-11700.000	5200.000		1.543	-176. 721
27	- 9 100. 000	4500.000	-178. 642 -178. 540	1.658	-176. 984
28	-9100. 000 -9100. 000	7800.000	-178. 568	1.454	-177. 114
20 29			-177. 786 170. 500	1.419	-176. 367
	-7800. 000 -4500. 000	6500.000 (500.000	-178. 522	1.521	-177. 001
30	-6500. 000	6500. 000 E200. 000	-179. 693	1.327	-17B. 366
31	-9100. 000	5200. 000	-178. 121	1.419	-176. 703
32	-10400.000	3900.000	-176. 078	1. 543	-174. 535
33	-13000.000	5200.000	-179. 363	1. 465	-177. 898
34	-14300.000	3900, 000	-180. 811	1. 521	-179. 290
35	-7800. 000	2600.000	-182. 305	1. 579	-180. 726
36	-6500. 000	2600.000	-182. 679	1. 607	-181.072
37	-3900. 000	2600, 000	-184. 023	1. 676	-182. 347
38	-7800. 000	1300.000	-182. 939	1.772	-181. 167
39	-7800. 000	5200. 000	-179. 256	1. 416	-177. B40
40	-7800. 000	7800. 000	-177. 536	1. 497	-176. 039
41	-6500. 000	7800.000	-178. 730	1.274	-177, 456
42	-6500. 000	5200.000	-180. 678	1. 376	-179, 302
4 3	-10400, 000	9100,000	-177. 641	1.489	-176, 152
44	-11700.000	7800, 000	-178. 470	1. 581	-176. 889
45	-13000.000	6500. 000	-179. 210	1. 457	-177. 753
46	-11700.000	6500.000	-178. 736	1. 582	-177. 154
47	-10400.000	7800.000	-178. 203	1. 457	-176. 746
48	-16900, 000	5200,000	-180. 811	1. 780	-179. 031

49	-13000, 000	-2600, 000	-185. 451	2, 074	-183. 377
50	-13000, 000	0.000	-182. 927	2. 051	-180. 876
 51	-15600, 000	0.000	-182. 327	1. 770	-180. 557
52	-15600.000	2600.000	-182. 137	1. 690	-180.447
53	-14300, 000	-1300.000	-183. 443	1, 993	-181. 450
54	-13000.000	1300.000	-182.102	1. 724	-180. 378
55	-11700.000	3900.000	-179. 036	1. 683	-177.353
56	~13000, 000	3900.000	-180. 056	1. 552	-178. 504
57	-19500, 000	6500 . 000	-179. 56 8	1. 422	-178. 146
58	-15600, 000	7800, 000	-179. 243	1. 374	-177. 869
59	-13000.000	9100.000	-178. 969	1. 375	-177. 594
60	-19500.000	9100.000	-178.075	1.344	-176. 731
61	-18200, 000	7800, 000	-178. 797	1. 325	-177. 472
62	-15600,000	-11700.000	-190. 746	2. 520	-188. 226
63	-10400.000	-10400. 000	-190. 994	2. 652	-188. 342
64	- 9100. 000	- 9100. 000	-190. 158	2. 677	-187. 481
45	-11700.000	-6500, 000	-187. 876	2. 495	-185. 381
66	-11700.000	-5200, 000	-188. 417	2. 465	-185. 952
67	-10400. 000	-2600, 000	-186. 883	3. 350	-183. 533
88	~7800. 000	-2600, 000	-186. 9 19	2. 219	-184. 700
69	-5200. 000	2600. 000	-183. 176	1.621	-181. 555
70	-22100. 000	6500. 000	-179. 876	1.386	-17B. 4 90
71	-24700.000	3900. 000	-180. 780	1. 593	-179. 187
72	-22100.000	3900, 000	-180. 728	. 1. 523	-179. 205
73	-24700.000	1300.000	-181. 169	1. 699	-179. 470
74	-23400. 000	-1300.000	-182. 547	1. 978	-180. 569
75	-22100.000	0. 000	-182. 727	1. 686	-181.041
76	-22100.000	1300.000	-182. 061	1. 630	-180. 431
77	-19500.000	3900.000	-181. 392	1. 541	-179. 851
78 70	-18200.000	`2600.000	-182. 084	1. 569	-180. 515
79	-18200, 000	0.000	-182. 347	1. 765	-180. 582
80	-20800.000	-1300.000	-183. 293	1.841	-181. 452
8 1 8 2	-16900, 000 -19500, 000	-2600, 000 -6500, 000	-181. 625	2. 026	-179. 599
83	-20800, 000 -20800, 000	-9100, 000 -9100, 000	-183. 207	2. 371	-180. 836
84	-15600, 000	-7800, 000 -7800, 000	-184. 580	2. 517	-182.063
85	-14300.000	-6500.000 -6500.000	-186, 147 -186, 674	2. 339 2. 459	-183. 808
86	-16900.000	-6500. 000 -6500. 000	-183. 885	2. 459 2. 611	-184. 215
87	-15600.000	-5200. 000	-183. 950 -183. 950	2. 240	-181, 274 -181, 710
88	-18200.000	-5200.000	-182. 073	2. 554	-179. 519
89	-14300.000	-3900.000	-184. 786	3. 301	-177. 317 -181. 485
90	-19500.000	-2600.000	-182. 890	2. 436	-180. 454
91	-22100.000	-6500.000	-184. 401	2. 339	-182.062
92	-24700.000	-5200.000	-183, 648	2. 163	-181.485
93	-19500.000	-10400, 000	-185. 795	2. 563	-183. 232
94	-16900, 000	-9 100.000	-186. 793	3.009	-183. 784
95	-13000, 000	-6500, 000	-188. 285	2.414	-185. 871
96	-5200.000	3900, 000	-183. 135	1.513	-181.622
97	- 6500, 000	3900, 000	-182.007	1.466	-180. 541
98	-3900. 000	3900, 000	-183, 893	1. 575	-182. 318
99	-5200. 000	5200.000	-182.000	1.415	-180. 585
100	- 7800. 000	3900, 000	-180. 650	1.478	-179. 172
101	-14300.000	-7800 , 000	-187. 609	2. 668	-184.941
 102	-13000.000	-7800, 000	-189. 095	2. 512	-186. 583
103	-13000.000	-9100.000	-190.032	2. 499	-187. 533
					- · · ·

104	-11700, 000	-7800, 000	-189. 516	2. 594	-186, 922
105	-22100.000	-3700, 000	-184, 218	2.002	-182, 217
106	-24700.000	-2600, 000	182. 291	1. 945	-180. 346
107	-28600.000	-1300.000	-176. 897	1. 969	-174. 928
108	-29900.000	3900.000	-178. 323	1. 663	-176. 660
					· ·
109	-36400.000	3900.000	-171. 860	1.849	-170. 011
110	-32500.000	0. 000	-174. 660	1. 926	-172. 734
111	-31200.000	-3900.000	-175. 593	2. 242	-173. 351
112	-28600, 000	-7800. 000	-179. 205	2. 296	-176. 909
113	-28 600. 000	-5200, 000	-178. 320	2. 225	-176. 095
114	-31200, 000	-6 5 00. 000	-175. 238	2. 263	-172. 975
115	-22100.000	-10400.000	-182. 849	2. 534	-180. 315
116	-24700.000	9100.000	-179, 963	1.400	-178. 563
117	-29900, 000	7800.000	-179, 422	1. 576	-177. 847
118	-9100.000	-13000.000	-191. 397	3. 201	-188. 196
119	-11700.000	-13000.000	-191.148	2. 773	-188. 375
120	-11700.000	-10400.000	-190. 984	2. 629	-188. 355
121	-6500, 000	-7800. 000 -7800. 000	-190.002	2. 874	-187. 128
122	-1340. 000				
		128.000	-185.088	2. 024	-183. 064
123	-38932. 000	-4030. 000	-171. 475	2. 101	-169. 374
124	-41120.000	-8080.000	-172. 280	2. 169	-170. 111
125	-46540. 000	-7410. 000	-168. 052	2. 444	-165. 608
126	-4 6260. 000	-964. 000	-1 67. 801	2. 423	-165. 378
127	-3660B. 00 0	10776. 000	-174. B62	1.842	-173. 020
128	-40300, 000	5200. 000	-169. 238	1. 924	-167. 314
129	-42511.000	-15334.000	<i>-</i> 173. 713	3. 200	-170.513
130	-28600. 00 0	-14300, 000	-181, 290	2. 969	-178. 321
131	-11700.000	0. 000	-183. 093	1. 937	-181. 156
132	-10400.000	5200.000	-177. 896	1. 510	-176. 386
133	-9100.000	5200.000	-178.078	1. 419	-176. 660
134	-13000.000	5200.000	-179. 326	1. 465	-177. 861
135	-14300.000	5200, 000	-180. 156	1. 466	-178. 690
136	-15600.000	5200 . 000			
137			-180. 550	1.443	-179. 107
	-10400.000	1300.000	-182. 258	1.860	-180. 398
138	-14300.000	3900. 000	-180. B75	1. 521	-179. 354
139	-5312, 000	0.000	-184. 782	1.859	-182. 923
140	-3660, 000	1300.000	-183. 852	1. 778	-182. 074
141	-4604. 000	-1117.000	-185. 042	2. 046	-182. 996
142	-6223. 000	-2454. 000	-186. 722	2. 184	-184. 538
143	-3084.000	-2704. 000	-185. 317	2. 772	-182. 545
144	-31200.000	-9 100.000	-176. 413	2. 336	-174. 077
145	-35655. 000	- 9696. 000	-174. 111	2.198	-171. 913
146	-26000.000	-11700.000	-182, 216	2. 607	-179, 609
147	-27300.000	-11700.000	-182. 195	2, 592	-179. 603
148	-24700.000	-13000.000	-181. 517	2. 754	-178. 763
149	-24700, 000	-14300.000	-182. 511	2. 830	-179. 681
150	-32177.000	-11879.000	-178. 603	2. 400	-176. 203
151	-32262.000	6336.000	-176. 516	1. 608	
152	-34355. 000	-156B. 000	-173. B99		-174. 908
152				2. 050	-171.849
ک.ن ∡	-18886. 000	-15115 000	-185. 424	2. B61	-182 563

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- D-3 Refraction and Resistivity Surveys
- D-3.1 Introduction

D-3.1.1 Purpose of Survey

Geophysical techniques were used in support of the subsurface hydrological investigation at the U.S. Army Depot near Tooele, Utah. The purpose of the investigation was to determine whether ground water contamination has occurred, and the potential for subsurface migration of contaminants. There were three objectives of the seismic refraction/electrical resistivity survey.

- 1. Determine subsurface structure south of the northern rock outcrop.
- 2. Determine if geophysical investigations can be used in the same area of TEAD to gather hydrologic information.
- 3. Provide a constraint for the interpretation of a gravity survey.

D-3.1.2 Survey Techniques

Three geophysical techniques; gravity, seismic refraction, and electrical resistivity, were used to delineate the subsurface structural features which may affect ground-water movement. A generalized basement-rock model of the valley was derived from a gravity survey conducted at the site. A description of the gravity survey can be found elsewhere in this report.

A seismic refraction survey was completed in the vicinity of the northern rock outcrop on the east side of the depot. The results of this survey were used to calibrate the gravity basement model and to confirm the existence of a shallow bedrock ridge that extends south from the outcrop. Such a ridge could restrict or alter the southeasterly flow of ground-water that has been inferred for the regional hydrogeologic system.



The refraction technique can delineate subsurface structure where an appropriate seismic velocity contrast exists. The technique measures the time required for a seismic wave to travel from a point of generation through the ground to detectors located on the surface. Wave arrival times are used to calculate the seismic velocities in the various media. Geologic stratigraphy and structure can be inferred by analyzing these velocities.

Electrical resistivity soundings were conducted at various locations along the refraction lines. This technique can delineate strata which have a resistivity contrast with the surrounding material. The method has been frequently useful in mapping ground-water aquifers because concentration and mobility of charged particles are the primary factors determining a material's resistivity.

A resistivity measurement is made in the following manner. A known current is forced to flow between two current electrodes which are in contact with the ground. This establishes an electric field in the earth. The distribution of the field depends on the resistivities and thicknesses of the various subsurface layers. The electric field is observed on the surface by measuring the potential difference across two potential electrodes. For this study, the potential electrodes are placed between, and in-line with, the current electrodes. One-dimensional, cross sections showing the distribution of resistivity layers with depth can be inferred from the measurements.

Significant ground-water in alluvium can usually be observed by both the seismic refraction and electrical resistivity techniques. The seismic compressional wave velocity in fine-grained materials such as sand or clay is

generally less than about 3000 feet/second (fps) (915 meters/second [mps]) when it is dry, but the velocity is between 4800 and 5500 fps (1463-1676 mps) when it is saturated. Saturated zones interpreted from seismic results should be correlated with other data because several dry earth materials also have characteristic velocities within this range.

Ground-water is usually slightly saline and normally appears as a low resistivity zone in the sounding results. A subsurface zone with a seismic velocity between 4800 and 5500 fps (1463-1676 mps) and a low resistivity value could be logically interpreted as being a saturated zone.

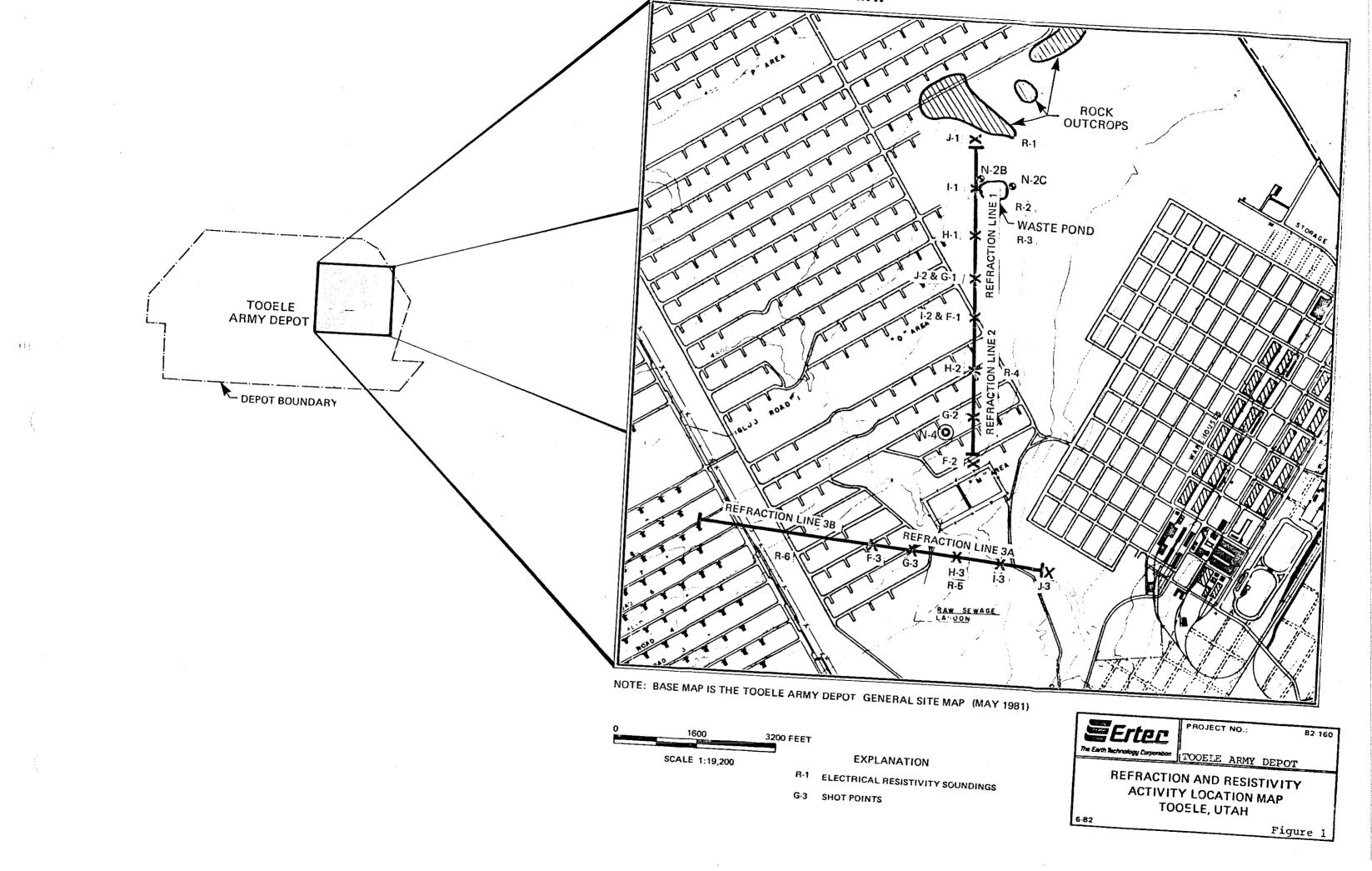
D-3.1.3 Scope of Work

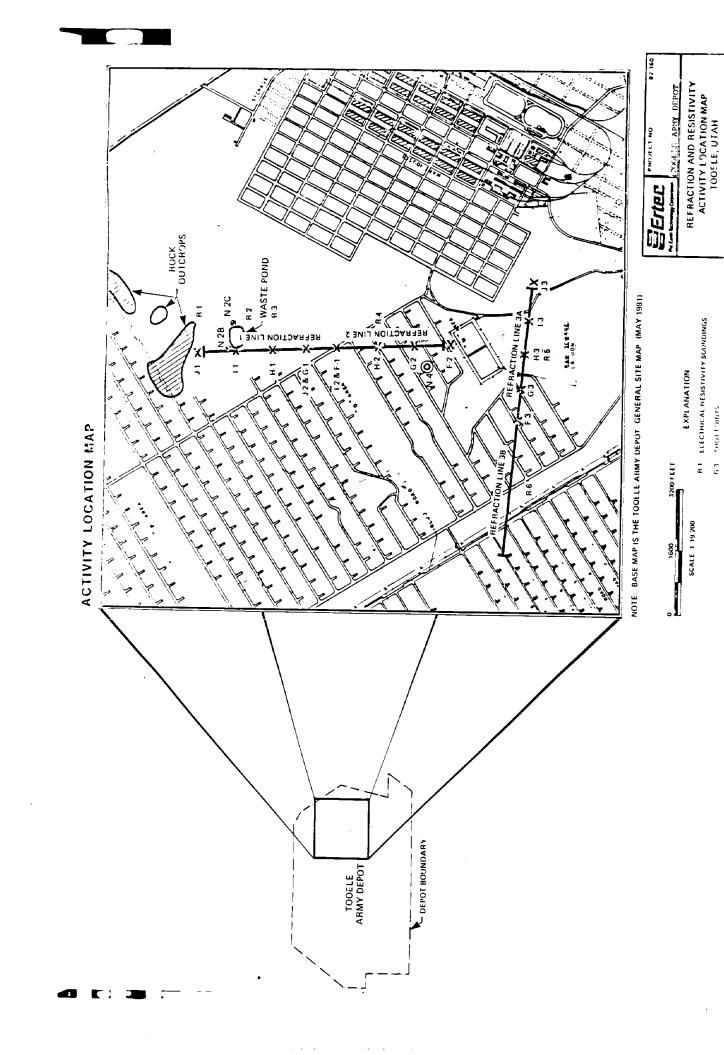
The seismic refraction survey is composed of four long lines as shown in Figure 1. The electrical resistivity survey consisted of six soundings which were conducted along the refraction lines. The survey was designed to collect information concerning the following questions:

- 1. Is the outcrop rock an isolated body or part of a more massive rock block?
- 2. If the outcrop is part of a larger rock mass, what is the depth and shape of the bedrock surface?
- 3. What is the velocity layering sequence in the overburden material?
- 4. Are there any geologic structures or unconformities in the section?
- 5. Are water saturated zones indicated?

The two north-south lines were positioned to map the buried rock ridge and the overburden material and the two east-west lines were positioned to investigate the lateral continuity of the section.







The desired depth of penetration of a refraction survey dictates the geophone spacing and the type of energy source used. The refraction lines were designed to penetrate about 600 feet, which was the approximate depths to bedrock estimated from examining existing wells logs and outcrops. Explosives were used to generate the seismic energy.

D-3.2 Seismic Refraction Survey

D-3.2.1 Recording Instruments

Seismic data were recorded with a 24-channel SIE Model RS-44 seismograph. The system processes 24 channels of seismic signals and produces a hard copy with a dry-write, galvanometer type recording oscillograph. The degree of gain was set on the system amplifiers by the instrument operator and was limited by the background noise at the time of the shot. Full width timing lines are impressed onto the record at 10 millisecond intervals. The timing lines are used to measure the time taken by the seismic wave to travel from the point of generation (shot point) to the geophones.

Mark Products Model L-10 geophones with a natural frequency of 4.5 Hz were used. They were fitted with short spikes to provide good coupling with the ground. Cables with connectors at 150-foot (46 m) intervals were used to transmit the signal from the geophones to the recording system.

D-3.2.2 Seismic Energy Generation

Seismic energy was generated by an explosive impulse. Small charges were placed in shallow borings at shot points along the refraction line. The charges were detonated one at a time and the resulting shock wave was measured

by surface detectors as it traveled through the ground. The safe and controlled use of explosives is essential in gathering refraction data. The U.S. Army required Ertec Western to demonstrate that proper safety procedures would be followed when handling explosive materials and to estimate the effect of vibrations produced by the explosions which would be used in the refraction survey.

In satisfying these requirements, the U.S. Army approved Ertec Western's "Standing Operating Procedures" for explosive use, then we conducted a blast vibration monitor program at the depot. A report (Ertec Western, 1982) was submitted describing the testing procedures and the recorded vibration levels. The approved procedures were followed during the refraction survey.

D-3.2.3 Refraction Data Acquisition

The field work was performed between January 8 and February 10, 1982. Each of the four refraction lines were about 3450 feet (1052 m) long and contained 24 geophones and five shot points as shown in Figure 1.

The geophone intervals and shot point locations were determined by the location of the connectors on the cables. Relative elevations of the geophones and shot points were measured with a transit and tied to true elevations on the existing grid at the depot. Geophones were placed and covered in shallow (less than 1 foot [0.3 m]) hand-excavated holes in order to improve energy coupling with the ground and to reduce extraneous wind noise. This procedure permitted the instrument operator to increase the amplification level in the recording system.

All shot points on a refraction line were drilled, loaded with explosives and backfilled before the recording began. One shot was denotated at a time and a



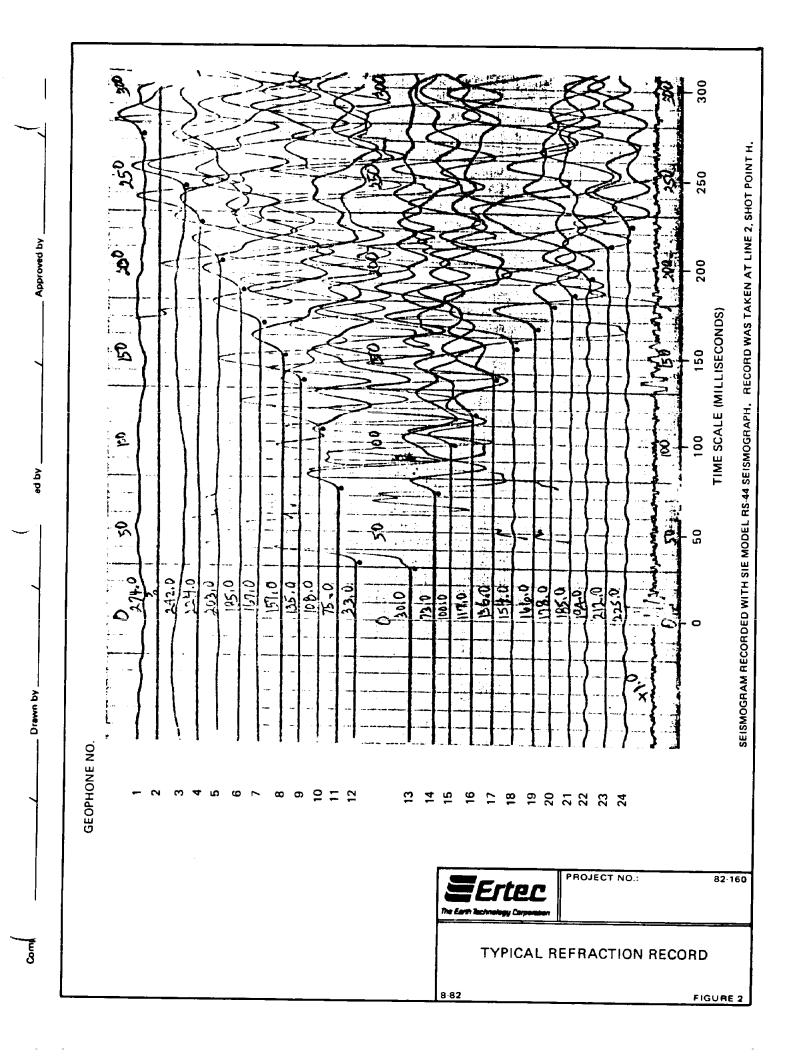
record was made of the event. The time of shot was transmitted from the shot point to the recording system via an FM radio link.

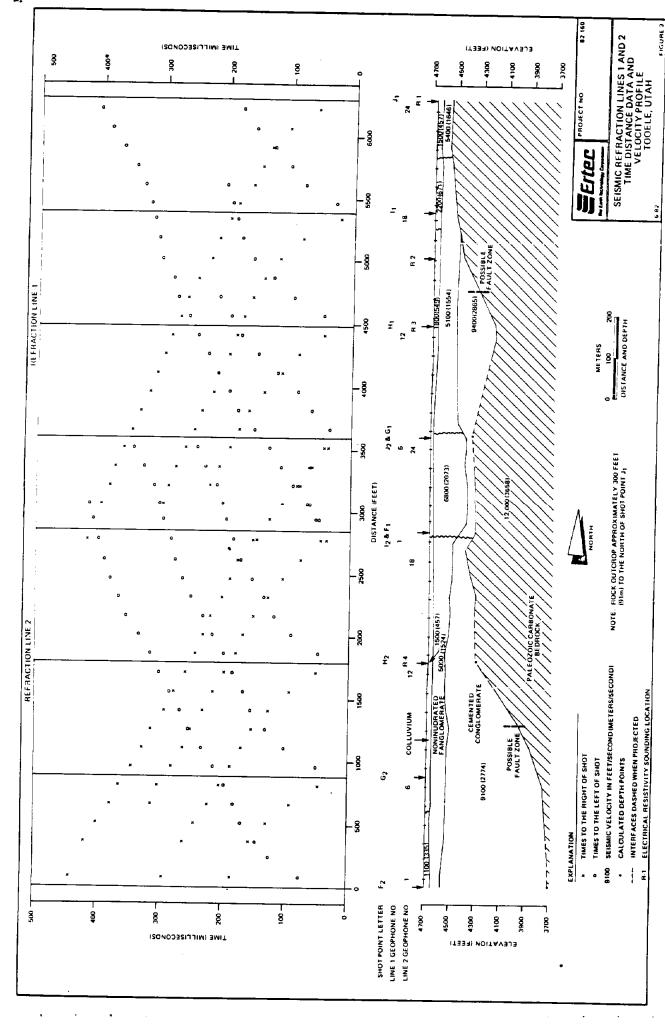
D-3.2.4 Refraction Data Reduction

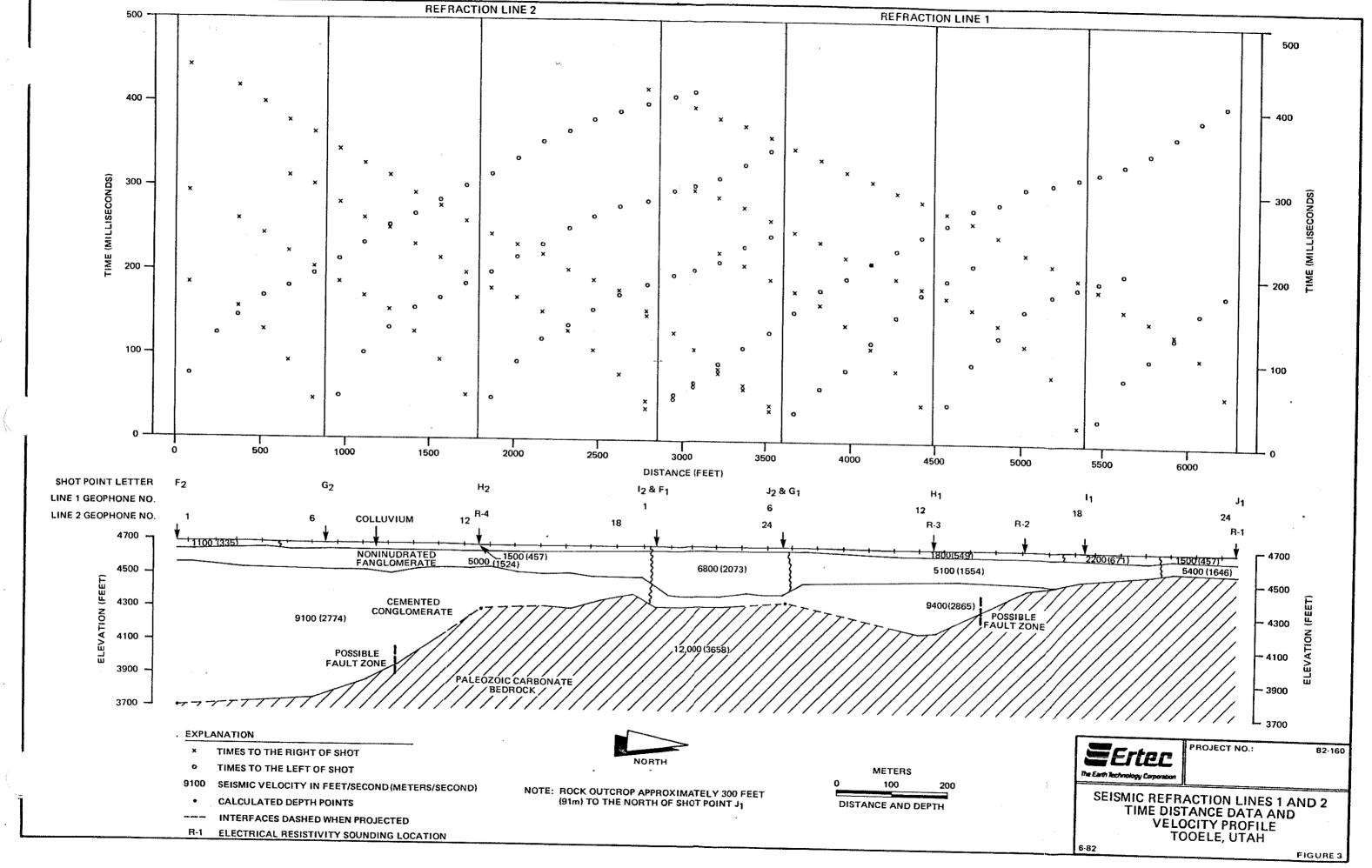
The refraction records were analyzed to determine the time interval between the explosive detonation and the arrival of the compressional (P) wave at each geophone. A typical refraction seismogram is shown in Figure 2. The P wave arrival usually appeared as a rather obvious excursion of the recording trace from its rest position. First, the compressional wave arrival at each geophone was identified on the record and then the travel times were measured. These times and the array geometry for each line are shown in Figures 3 and 4.

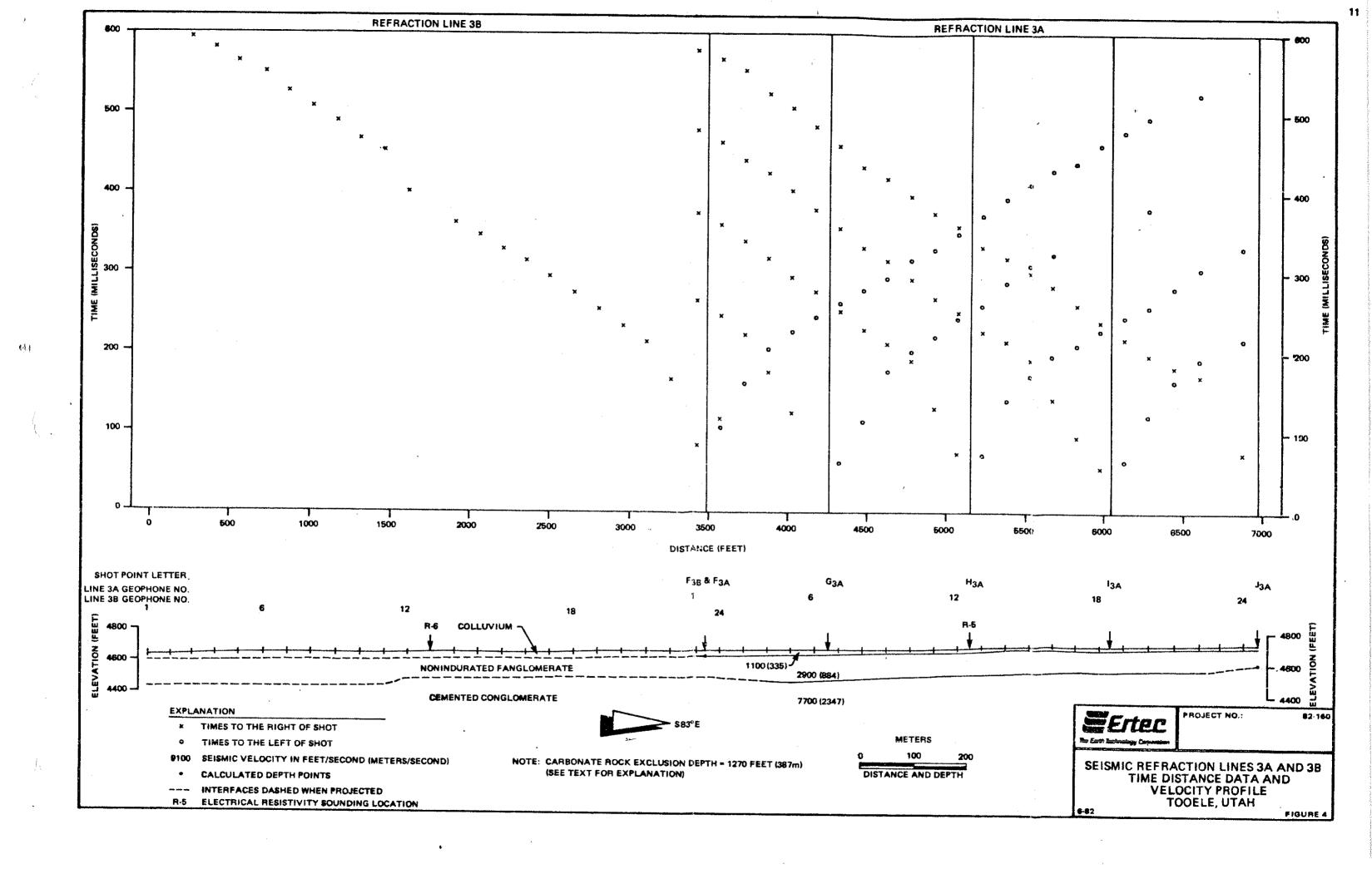
The upper halves of these figures are graphs of seismic wave travel times plotted versus surface distance between the shot points and the geophones. Distances were measured along the north-south line from geophone number 1, of line 2 and along the east-west line from geophone 1 of line 3B. Distances to the right of the first geophone are positive. The direction arrow gives the approximate orientation of the line from the first geophone toward the other end of the line. The vertical lines designate the locations of shot points (F, G, H, I, J). The symbol X denotes travel times at geophones that were to the right of a shot point. The symbol 0 denotes travel times to geophones that were to the left of a shot point.

The Generalized Reciprocal Method (GRM) of seismic refraction interpretation (Palmer, 1980) was used to analyze these graphs and to calculate the velocity profile beneath each line. Computer programs SEISSF and DSECTN (Hatherly, 1978) were used to calculate the various functions of the GRM. The GRM combines the computational ease of the reciprocal method (Hawkins, 1961) with the migration property of the delay time method (Gardner, 1939).









The lower halves of Figures 3 and 4 show the interpreted seismic velocity profiles. The top lines represent the ground-surface profiles. The short vertical lines crossing the top line mark the geophone positions. The depth scales are plotted relative to true elevations from sea level. The additional lines across the cross section represent the interpreted boundaries between layers of material with different compressional wave velocities. The calculated P wave velocity is shown for each layer.

D-3.2.5 Refraction Results

A brief description of the refraction results for the north-south and eastwest lines is given below. The correlation between velocities and lithologies is based on interpretations of boring logs obtained during the hydrologic investigation.

North-South Line (Figure 3)

- 1. The first layer (surface material) is a colluvium that has a very low average P wave velocity of 1600 fps (488 mps). The thickness of the layer varies between 25 and 50 feet (8 and 15 m). The measured velocities for this layer varied by 31 percent from 1100 to 2200 fps (335 to 671 mps).
- 2. The second layer is a nonindurated fanglomerate that has an average P wave velocity of 5150 fps (1570 mps). The velocity of this layer varies by 17 percent except for an anomaly, 6800 fps (2073 mps), in the center of the line. The top surface of the layer is fairly flat except near the velocity anomaly. The thickness ranges between 70 and 270 feet (21 and 82 m).
- 3. The third layer in the section is a cemented conglomerate that has a well defined, average P wave velocity of 9250 fps (2819 mps) which varied by \pm 2 percent. This layer is 850 feet (259 m) thick on the south end of the line and pinches out just south of shot point I_1 . The geometry of this layer is irregular.
- 4. The deepest velocity layer mapped by this line is a Paleozoic carbonate bedrock with a P wave velocity of 12,000 fps (3658 mps) which varied by ± 8 percent. The top surface of this layer is irregular.

East-West Line (Figure 4)

- 1. The colluvium, surface layer has a fairly well defined P wave velocity of 1100 fps (335 mps). This velocity varied + 18 percent from 950 to 1300 fps (290 to 396 mps). The layer has an average thickness of 25 feet (8 m).
- 2. The second layer, fanglomerate, has an average P wave velocity of 2900 fps (884 mps) that varied by + 10 percent. The surface of this layer is fairly flat with thicknesses ranging from 130 to 210 feet (40 to 64 m).
- 3. The deepest velocity layer mapped under this line is a cemented conglomerate that has an average P wave velocity of 7700 fps (2347 mps) that varied by ± 9 percent. The top surface of this layer is fairly flat. There is a 40-foot (12 m) step, in the calculated layer surface at line 3B, geophone 11.
- 4. The 12,000 fps (3658 mps) velocity associated with carbonate bedrock was not observed with the east-west lines. Therefore, the bedrock must be beyond the depth of penetration associated with the line geometry. This geometry would have detected the 12,000 fps (3658 mps) layer if it had been shallower than 1270 feet (387 m).

The P wave arrivals on the refraction records are good quality and are easily identified. The seismic velocities are estimated to be correct within \pm 10 percent or less, except for the surface layer. Arrivals representing seismic waves traveling solely in the first layer are seldom observed at more than one geophone from any shot. Therefore, the accuracy of the first layer velocity determination is relatively low because they are based on so few data points.

D-3.3 ELECTRICAL RESISTIVITY SURVEY

D-3.3.1 Recording Instrument

The electrical resistivity data were measured with a Bison Model 2350A resistivity meter. Stainless steel rods were used for the current electrodes. Copper rods immersed in a saturated copper sulfate solution in a porous pot were used for the voltage measuring electrodes.



D-3.3.2 Resistivity Data Acquisition

The field data were collected between January 28 and February 18, 1982. Six soundings using the Schlumberaer electrode array were made. Their locations are shown in Figure 1. The geometry of this measurment array is shown in Figure 5. To make a sounding, the spacing between the electrodes is expanded between measurements, while the center of the array remains fixed. As the array expands, the current penetrates deeper into the earth, thus allowing the variation of resistivity with depth and material changes to be reflected in the surface measurements. Fourteen measurements were taken for each sounding, with the electrode separation (Figure 5) expanding from 15 to 350 feet (5 to 107 m).

Apparent resistivity values were calculated from the measured values using the following equation:

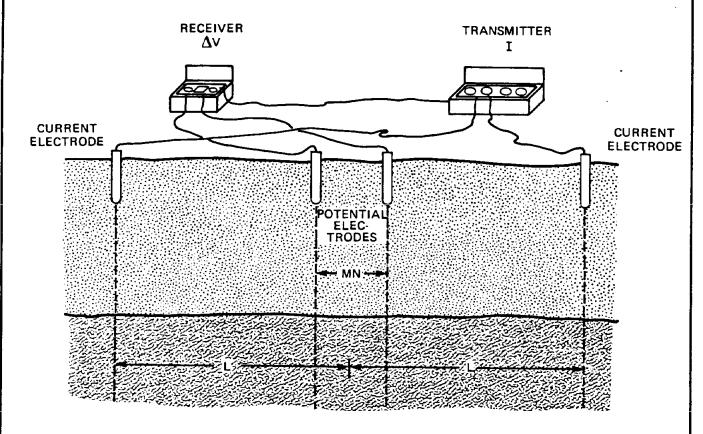
$$= \frac{L^2}{MN} \frac{V}{I} (ohm meter)$$

(See Figure 5 for definition of the various terms). The apparent resistivity was plotted versus the electrode separation and checked for bad points by the field crew before leaving the test site. The resistivity data appears in the upper half of Figures 6 through 11.

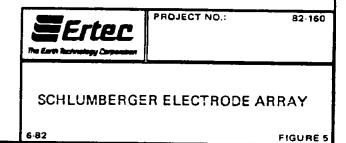
D-3.3.3 Resistivity Data Reduction

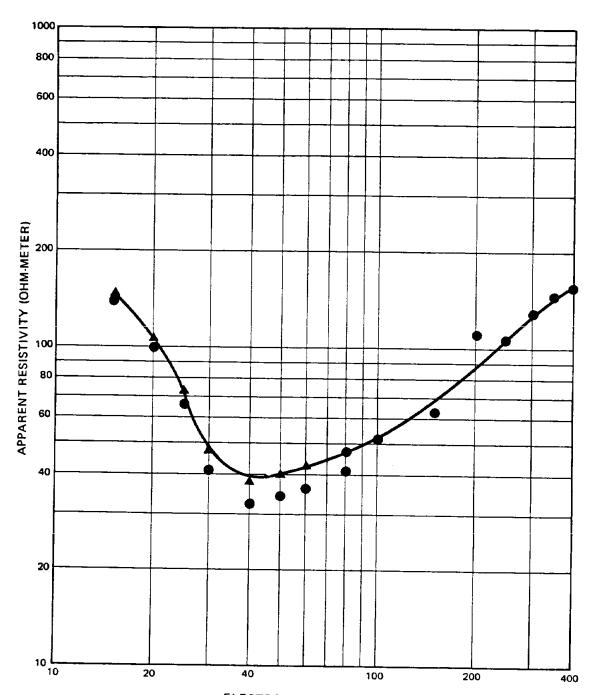
If the ground is electrically homogeneous, the measured resistivity values would represent the true soil resistivities. However, the ground is usually composed of several subsurface layers, each of which will alter the distribution of the applied electric field. Therefore, the surface measurements represent a composite resistivity value of the underlying strata. This composite resistivity is termed "apparent resistivity" and is not equal to true





SCHLUMBERGER ARRAY





● OBSERVED VALUES
▲ ADJUSTED VALUES

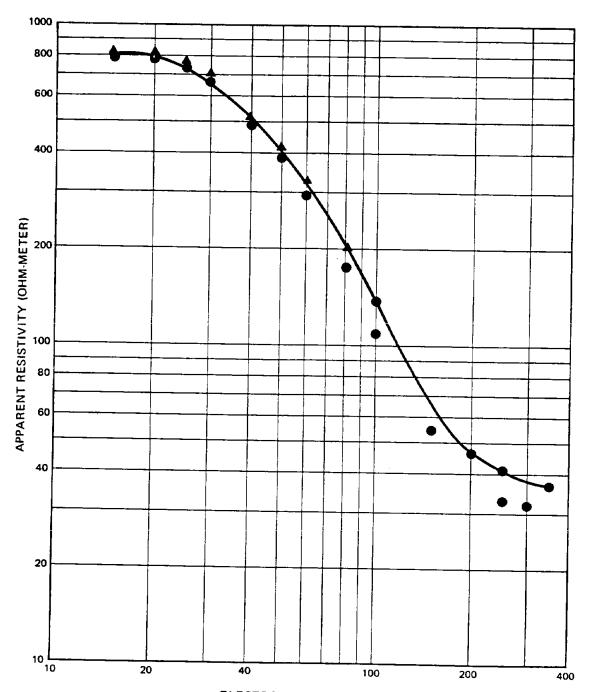
ELECTRODE SPACING - AB/2 (FEET)

6-82

		INTERPRET	TED MODEL	
LAYER DEPTH RESISTIVITY VALUES				TY VALUES
FEET	METERS	% DEVIATION	OHM-METER	% DEVIATION
0	0		310	11
7	2	5	30	5
59	18	11	460	28

Ertec	PROJECT NO.:	82-160
The Earth Technology Corporation		

RESISTIVITY SOUNDING R-1 SOUNDING CURVE AND INTERPRETATION TOOELE, UTAH



OBSERVED VALUES

ELECTRODE SPACING - AB/2 (FEET)

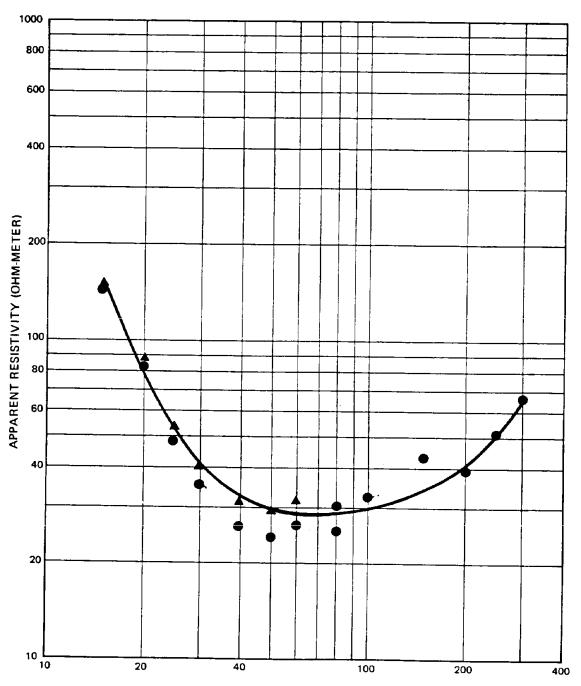
▲ ADJUSTED VALUES

		INTERPRET	TED MODEL	
	LAYER	DEPTH	RESISTIVI	TY VALUES
FEET	METERS	% DEVIATION	OHM-METER	% DEVIATION
0	0	_	720	4
31	9	3	40	5
	<u> </u>			

	,	
Ertec	PROJECT NO.:	82-160
The Earth Rechnology Corporation		

RESISTIVITY SOUNDING R-2 SOUNDING CURVE AND INTERPRETATION TOOELE, UTAH

6.82



▲ ADJUSTED VALUES

ELECTRODE SPACING - AB/2 (FEET)

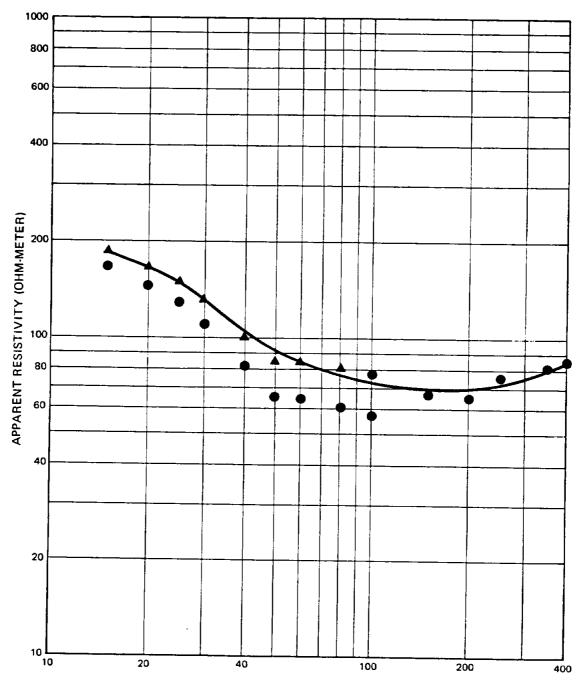
		INTERPRET	ED MODEL	
	LAYER	DEPTH	RESISTIVI	TY VALUES
FEET	METERS	% DEVIATION	OHM-METER	% DEVIATION
0	0		230	18
7	2	9	30	7
ຍງ	27	39	130	94

(Table)	PR
Erter	!
	l
The Earth Richnology Corporation	

ROJECT NO.: 82-160

RESISTIVITY SOUNDING R-3 SOUNDING CURVE AND INTERPRETATION TOOELE, UTAH

6.82



ELECTRODE SPACING - AB/2 (FEET)

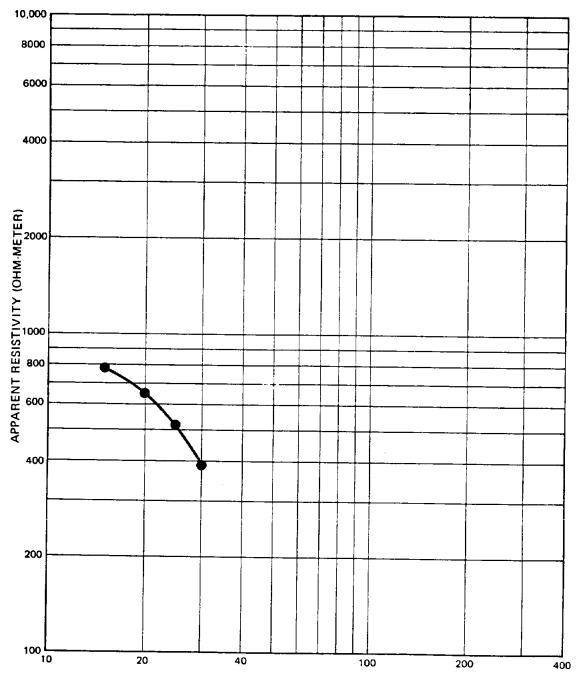
▲ ADJUSTED VALUES

		INTERPRET	TED MODEL	
	LAYER	DEPTH	RESISTIVI	TY VALUES
FEET	METERS	% DEVIATION	OHM-METER	% DEVIATION
0	0	-	210	7
13	4	8	70	12

Ertec	PROJECT NO.:	82-160
The Earth Richnology Corporation	<u> </u>	

RESISTIVITY SOUNDING R-4 SOUNDING CURVE AND INTERPRETATION TOOELE, UTAH

6.8



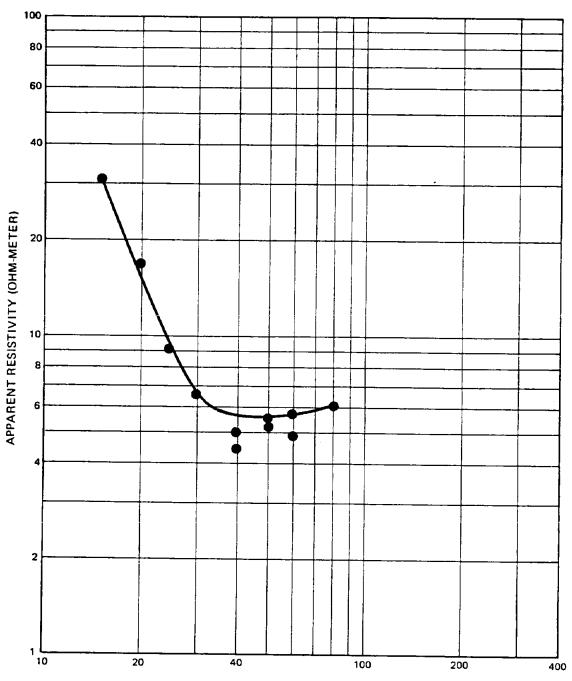
ELECTRODE SPACING - AB/2 (FEET)

INTERPRETED MODEL					
LAYER DEPTH RESISTIVITY VALUES				TY VALUES	
FEET	METERS	% DEVIATION	OHM-METER	% DEVIATION	
		NOT ENOUGH D	ATA TO MODEL		
L					

E rtec	PROJECT NO.:	82-16

RESISTIVITY SOUNDING R-5 SOUNDING CURVE AND INTERPRETATION TOOELE, UTAH

6.82



ELECTRODE SPACING - AB/2 (FEET)

	-	INTERPRET	TED MODEL	· · · · · · · · · · · · · · · · · · ·	
LAYER DEPTH RESISTIVITY VALUES					
FEET	METERS	% DEVIATION	OHM-METER	% DEVIATION	
0	0	-	110	6	
6	_ 2	2	5	3	
42	13	72	9	67	

EFTEC	PROJECT NO.:	82-160
	·	

RESISTIVITY SOUNDING R-6 SOUNDING CURVE AND INTERPRETATION TOOELE, UTAH

6-82 FIGURE 11

soil resistivity. By making a number of measurements at a site, it is possible to convert an apparent resistivity field curve into a layered subsurface model with a calculated "true" resistivity for each layer of the model.

A resistivity field curve consists of several segments, each corresponding to a differend MN spacing. the vertical offset between two adjacent segments is caused by either one or both of the potential electrodes crossing a near surface, lateral interface into a material with a different resistivity value. These offsets must be removed prior to computer modeling because they represent a resistivity change due to lateral not depth effects. The segments were adjusted relative to the largest MN spacing used in the sounding. At this spacing, a large part of the electric field in the earth is being measured. This large measurement interval is not an influenced by small, localized, surface resistivity anomolies as smaller MN spacings. Consecutive segments (from large to small MN spacings) were vertically shifted until a smoothed curve was obtained. The curve was then digitized and used for computer input.

The resistivity data were interpreted with the aid of two inverse modeling computer programs. The program by Zhody (1973) uses modified Dar Zarrouk functions in an iterative process to determine true resistivity values of the various subsurface layers. The resulting model is then used as an initial estimate to Rijo's (1977) program. The program calculates theoretical resistivity values by convolution with a set of filter coefficients. Derivatives of the theoretical resistivity values with respect to the parameters are then calculated numerically and used in a Marquardt (ridge) regression algorithm to calculate improvements in the values of the parameters. This process is continued through a specified number of iterations or until the successive changes in the parameters is less than a preselected value.

The resulting geoelectric cross sections were evaluated versus the field curves and geologic knowledge of the area. The interpreted models are tabulated at the bottom of Figures 6 through 11.

D-3.3.4 Resistivity Results

The interpretation of the electrical resistivity survey is shown in cross sectional view in Figure 12. The four resistivity soundings along the north-south refraction line show the top of a low resistivity zone between 10 and 30 feet (3 and 9 m) deep. This low resistivity zone is in the fanglomerate material. Soundings R-1 and R-3 show this zone to be fairly thick, about 60 feet (18 m). The seismic velocity of this layer is between 5000 and 5400 fps (1524 and 1646 mps). Therefore this layer could be interpreted as being a saturated zone of varying thickness.

Sounding R-6 shows a thick, low resistivity layer near the surface which also corresponds with the fanglomerate layer. The seismic velocity of this layer is 2900 fps (884 mps) which is well below the characteristic value for water. Therefore, if any saturated zones exist under the east-west refraction line, they must be small and isolated.

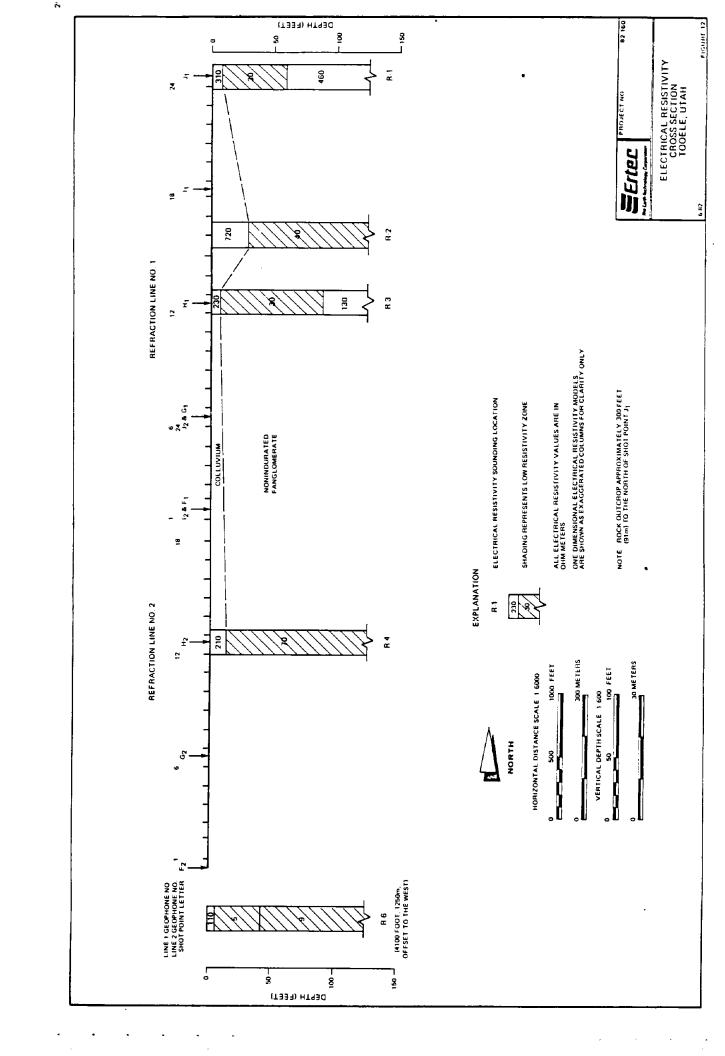
D-4 INTEGRATION OF RESULTS

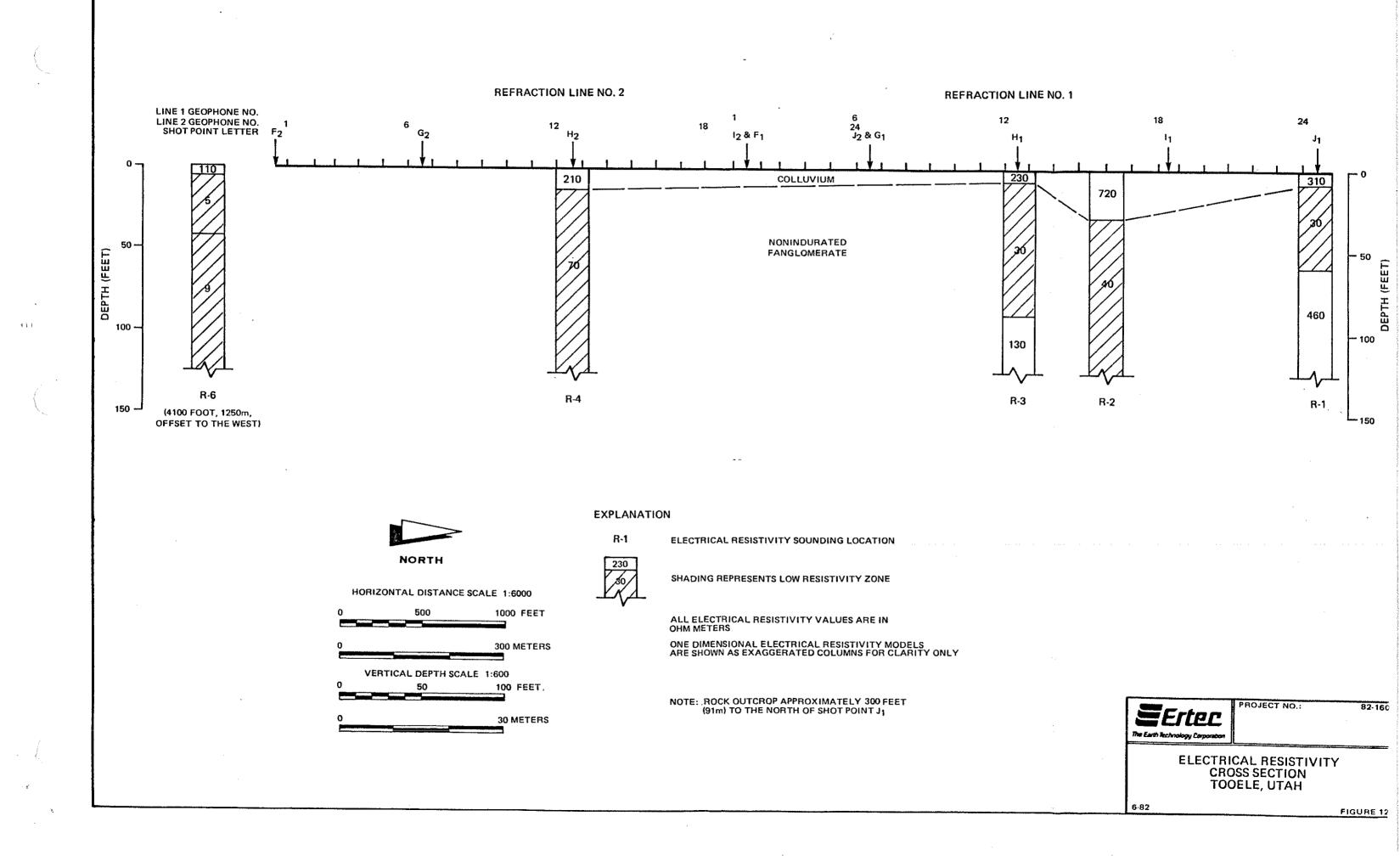
The refraction/resistivity surveys have indicated answers to the questions posed in D-3.1.3.

1. Nature of Outcrop

The northern rock outcrop is the surface expression of a much larger rock mass. The measured seismic velocity of the rock is 12,000 fps (3658 mps), which is within the range of velocities typically measured in carbonate rock.







2. Bedrock Surface

The rock surface dips towards the south in a series of terraces. The three terraces are probably separated by two fault zones (Figure 3). The average depths of the terraces beneath the seismic line are:

- o 180 feet (55 m) northern terrace
- o 400 feet (122 m) central terrace
- o 950 feet (290 m) southern terrace

The carbonate rock is not detected by the east-west refraction line which is 1800 feet (549 m) south of the north-south line. The depth to which we could exclude the rock under this line was calculated to be 1270 feet (387 m).

3. Overburden Layering

There are three distinct subsurface velocity layers overlying the bedrock. They correlate with:

Colluvium -

mostly unconsolidated sands, silts and gravel.

Non-indurated fanglomerate - a variable-cobble matrix filled with unconsolidated, fine-grained material.

Cemented conglomerate -

a variable-cobble matrix filled with cemented finegrained material.

The inhomogeneity in the colluvium is expressed in the wide range of observed velocities, 1100 to 2200 fps (335 to 671 mps). The zone of non-indurated fanglomerate has a velocity ranging from about 5100 fps (1554 mps) on the N-S line to 2900 fps (884 mps) on the E-W line. The cemented conglomerate also has a higher velocity (9100 to 9400 fps; 2774-2865 mps) beneath the N-S line than it does beneath the E-W line, 7700 fps (2347 mps). The differences in the velocity layering between the two lines may be caused by anisotropy, differences in the materials, differences in the moisture content, or a combination of these factors.

4. Geologic Structures

Several structural features are interpreted from the refraction survey. Two bedrock fault zones are interpreted beneath the north-south refraction lines. They are located approximately 2000 and 6000 feet (610 and 1829 m) south of the outcrop. The approximate vertical offsets on the faults are:



150 feet (46 m) northern fault zone 550 feet (168 m) southern fault zone

The east-west line was positioned to cross a large gravity gradient which might indicate a fault. The seismic data indicates a vertical offset in the 7700 fps layer near the center of line 3B, which is also near the gravity gradient. The offset in this layer is on the order of 40 feet (12 m) but it could be caused by a larger displacement in the bedrock. Bedrock structure at this location is too deep to be determined by the refraction line geometry.

Another structure is indicated by the velocity anomaly in the fanglomerate material at the center of the north-south line. The velocity of the anomaly is 6800 fps (2073 mps) which is about 33 percent higher than the velocity of the surrounding material. The cause of the anomaly is interpreted to be a buried stream channel. The base of the channel is about 700 feet (213 m) wide and about 300 feet (91 m) below the surface. The higher velocity may indicate that material in the channel is much coarser than the surrounding material.

5. Water Saturated Zones

The results of the two surveys indicate that there may be a perched water zone beneath the north-south line, but no significant ground water under the east-west line. The perched water zone may exist at the north-south line because of well-developed cementation of the underlying conglomerate. The fact that the velocity in this zone is higher than it is beneath the east-west line indicates better cementation beneath the north-south line.

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D-5 Horizontal and Vertical Control Results

TOOELE ARMY DEPOT / NORTH LAMBERT - UTAH CENTRAL

		LONGITUDE	NORTHING EASTING
4 0 - 0	32 35.292 33 06.090 4679.190	112 21 40.463 CONVERGENCE ELEVATION	806,047.120 1,760,638.300 0.9999752 SCALE FACTOR 0.9997870 GRID FACTOR
- 40	33 04.500	112 21 37.981 CONVERGENCE ELEVATION	805,581.300 1,760,825.480 0.9999749 SCALE FACTOR 0.9997866 GPID FACTOR
- 40	34 27.925	112 23 48.214 CONVERGENCE ELEVATION	792,717.360 1,750,641.410 0.9999674 SCALE FACTOR 0.9997774 GRID FACTOR
40	34 27.843	112 23 48.087 CONVERGENCE ELEVATION	792,714.920 1,750,651.210 0.9999674 SCALE FACTOR 0.9997774 GFID FACTOR
- 40	33 15.340	112 21 54.903 CONVERGENCE ELEVATION	801,178.630 1,759,476.340 0.9999723 SCALE FACTOR 0.9997847 GPID FACTOR
- 0	30 03.267 36 57.525 5094.340	112 27 41.754 CONVERGENCE ELEVATION	790,948.050 1,732,581.600 0.9999663 SCALE FACTOR 0.9997616 GFID FACTOR
- 0	33 26.853	112 22 12.876 CONVERGENCE ELEVATION	787,720.530 1,757,957.540 0.9999647 SCALE FACTOR 0.9997696 GRID FACTOR
- 0	33. 50.598 34 11.342 4474.530	112 23 22.327 CONVERGENCE ELEVATION	813,744.460 1,752,850.250 0.9999798 SCALE FACTOR 0.9997997 GFID FACTOR
- 0	33 50.598 34 11.482 4474.940	112 23 22.545 CONVERGENCE ELEVATION	813,744.610 1,752,833.470 0.9999798 SCALE FACTOR 0.9997997 GFID FACTOR

TOOELE ARMY DEPOT / NORTH U.T.M.- ZONE # 12

LATITUDE	LONGITUDE	NOFTHING	EASTING
40 32 35.292 0 53 05.747 4679.190	112 21 40.463 CONVERGENCE ELEVATION	4,488,722.030 0.9997635 SCALE FACTOR 0.9995398 GPID FACTOR	384,726.049
0 53 04.050	112 21 37.961 CONVERGENCE ELEVATION	4,488,579.753 0.9997633 SCALE FACTOR 0.9995394 GFID FACTOR	384,782.252
0 54 26.357	112 23 48.214 CONVERGENCE ELEVATION	4,484,677.706 0.9997723 SCALE FACTOR 0.9995463 GRID FACTOR	381,656.041
0 54 26.274	112 23 48.087 CONVERGENCE ELEVATION	4,484,676.950 0.9997723 SCALE FACTOR 0.9995463 GFID FACTOR	381,659.019
0 53 14.263	112 21 54.903 CONVERGENCE ELEVATION	4,487,240.514 0.9997645 SCALE FACTOR 0.9995415 GFID FACTOR	384,363.336
0 56 57,728	112 27 41.754 CONVERGENCE ELEVATION	4,484,170.537 0.9997887 SCALE FACTOR 0.9995452 GFID FACTOR	376,149.382
0 53 23.526	112 22 12.876 CONVERGENCE ELEVATION	4,483,142.053 0.9997659 SCALE FACTOR 0.9995338 GRID FACTOR	383,876.674
40 33 50.598 0 54 13.370 4474.530	112 23 22.327 CONVERGENCE ELEVATION	4,491,081.449 0.9997703 SCALE FACTOR 0.9995563 GPID FACTOR	382,366.442
40 33 50.598 0 54 13.512 4474.940	112 23 22.545 CONVERGENCE ELEVATION	4,491,081.530 0.9997703 SCALE FACTOR 0.9995563 GRID FACTOR	382,361.315

IOOELE ARMY DEPOT / NORTH LAMBERT - UTAE CENTFAL

	LATITUDE	LONCITUDE	NCFTHINC EASTING	SAMPLE #
4 (CONTRACTOR	792,581.500 1,750,701.400 0.9999673 SCALE FACTOR 0.9999673 GFID FACTOR	
4 0 - 0	33 05.998	112 21 40.319 CONVERGENCE ELEVATION	805,676.900 1,760,645.900 0.9999749 SCALE FACTOR 0.9999749 GFID FACTOR	NSW-2
- 40 - 0	33 12.033	112 21 49.740 CONVERGENCE ELEVATION	800,252.600 1,759,866.100 0.9999718 SCALE FACTOR 0.9999718 GFID FACTOR	NSW-3
40	35 58.307	112 26 09.309 CONVERGENCE ELEVATION	0-9999670 SCALE BACKOR	NSD-1
40 - 0	35 31.453	112 25 27.387 CONVERGENCE ELEVATION	813,583.200 1,743,196.000 0.9999796 SCALE FACTOR 0.9999796 GFID FACTOR	NSD-2
40 - 0	33 05.99E	112 21 40.319 CONVERGENCE ELEVATION	0.9999749 SCALE BACKER	NSD-3

TOOELE ARMY DEPOT / NORTH U.T.M.- ZONE # 12

	LATITUD	E LONGITUDE	NORTHING EASTING	- S SAMPLI
4 ((0 54 25.81	8 112 23 47.420 6 CONVERGENCE 0 ELEVATION	4,484,636.216 381,674.077 0.9997723 SCALE FACTOR 0.9997723 GRID FACTOR	NSW-1
4 0	53 05.58	4 112 21 40.319 7 CONVERGENCE 0 ELEVATION	4,488,609.186 384,727.694 0.9997635 SCALE FACTOR 0.9997635 GRID FACTOR	NSW-2
4 0 - 0	0 53 10 ₄ 74	0 112 21 49.740 2 CONVERGENCE 0 ELEVATION	4,486,957.610 384,480.466 0.9997642 SCALE FACTOF 0.9997642 GRID FACTOR	NSW-3
40 - 0	55 57.88	5 112 26 09.309 CONVERGENCE DELEVATION	0.9997821 SCALE FACTOR	NSD-1
40 · 0	55 34.676	3 112 25 27.387 CONVERGENCE ELEVATION	4,491,049.484 379,424.190 0.9997789 SCALE FACTOR 0.9997789 GFID FACTOR	NSD-2
4 0 0	53 05.587	112 21 40.319 CONVERGENCE ELEVATION	4,488,609.186 384,727.694 0.9997635 SCALE FACTOR 0.9997635 CLID FACTOR	NSD-3

TOOELE ARMY DEPOT / SOUTH LAMBERT - UTAH CENTRAL

		LONGITUDE	
40 17 2	1.978 112	2 19 22.208	713,531.220 1,770,461.39
0 31 3	7.527 CON	EVERGENCE	0.9999302 SCALE FACTOR
514	8.420 ELE	EVATION	0.9997239 GFID FACTOR
			716,593.320 1,756,677.58 0.9999313 SCALE FACTOR 0.9997288 GRID FACTOR
40 15 4 0 31 3	4.336 112 2.696 CON	2 19 14.667 VERGENCE	703,646.170 1,770,955.03 0.9999266 SCALE FACTOR 0.9997237 GRID FACTOR
40 15 4	3.909 112	19 42.724	703,623.070 1,768,779.69
0 31 5	0.669 CON	VERGENCE	0.9999266 SCALE FACTOR
505	2.070 ELE	VATION	0.9997242 GPID FACTOF
40 15 58	8.659 112	21 45.648	705,205.600 1,759,265.33
0 33 09	9.412 CON	VERGENCE	0.9999271 SCALE FACTOR
5039	9.930 ELE	VATION	0.9997252 GRID FACTOR
40 16 23	1.838 112	21 00.702	707,517.530 1,762,771.55
0 32 40	0.620 CON	VERGENCE	0.9999280 SCALE FACTOR
5048	8.770 ELES	VATION	0.9997257 GRID FACTOR
40 17 51	760 112	19 12.931	716,538.060 1,771,207.81
0 31 33	585 CONV	VERGENCE	0.9999314 SCALE FACTOR
5190	0.180 ELEV	VATION	0.9997234 GPID FACTOR
40 16 14	.999 112	19 17.013	706,750.400 1,770,801.65
0 31 34	.199 CONV	PERGENCE	0.9999277 SCALE FACTOR
5101	.630 ELEV	PATION	0.9997234 GRID FACTOR
40 16 52	.782 112	18 50.643	710,554.720 1,772,880.39
0 31 17	.307 CONV	ERGENCE	0.9999291 SCALE FACTOF
5125	.600 ELEV	ATION	0.9997238 GRID FACTOR

	30 32.605	112 17 40.858 CONVERGENCE	721,052.290 1,778,382.100 0.9999332 SCALE FACTOR	S-11
	5353.780	ELEVATION	0.9997186 GPID FACTOR	
		112 23 33.635	703,409.150 1,750,876.230	S-12
- 0		CONVERGENCE	0.9999264 SCALE FACTOR	_
	5054.360	ELEVATION	0.9997240 GPID FACTOR	
40	17 10.651	112 23 51.558	712,585.810 1,749,578,900	S-14
- 0	34 30.067	CONVERGENCE	0.9999298 SCALE FACTOR	3-14
	5039.130	ELEVATION	0.9997279 GFID FACTOR	
40	18 51.051	112 18 52.056	722,522.460 1,772,879.800	S-15
- 0	31 18.212	CONVERGENCE	0.9999337 SCALE FACTOR	J=1.)
		ELEVATION	0.9997207 GRID FACTOR	

TOOELE ARMY DEPOT / SOUTH
U.T.M.- ZONE # 12

	LONGITUDE	
40 17 21.978 0 51 19.796 5148.420	112 19 22.208 CONVERGENCE ELEVATION	4,460,511.704 387,556.718 0.9997556 SCALE FACTOR 0.9995094 GRID FACTOR
		4,461,468.967 383,361.543 0.9997674 SCALE FACTOR 0.9995258 GFID FACTOR
40 15 44.336 0 51 13.201	112 19 14.667 CONVERGENCE	4,457,498.459 387,689.907 0.9997552 SCALE FACTOR 0.9995130 GRID FACTOR
40 15 43.909 0 51 31.332 (5052.070 F	112 19 42.724 CONVERGENCE ELEVATION	4,457,495.198 387,026.955 0.9997570 SCALE FACTOR 0.9995155 GFID FACTOR
40 15 58.659 J 0 52 51.070 d	112 21 45.648 CONVERGENCE	4,457,994.065 384,130.261 0.9997652 SCALE FACTOR 0.9995242 GRID FACTOR
40 16 21.838 1 0 52 22.426 C 5048.770 E	12 21 00.702 CONVERGENCE LEVATION	4,458,692.494 385,202.791 0.9997622 SCALE FACTOR 0.9995208 GRID FACTOR
40 17 51.760 1 0 51 14.319 C	.12 19 12.931	4,461,426.705 387,789.456 0.9997549 SCALE FACTOR 0.9995068 GRID FACTOR
U 51 15.257 C	12 19 17.013 ONVERGENCE LEVATION	4,458,444.713 387,648.586 0.9997553 SCALE FACTOR 0.9995114 GPID FACTOR
U DU DB.868 CI	12 18 50.643 ONVEFGENCE LEVATION	4,459,600.412 388,288.688 0.9997536 SCALE FACTOR 0.9995085 GFID FACTOR

- 4	0 50 15.532	112 17 40.858 CONVERGENCE ELEVATION	4,462,769:841 0.9997489 SCALE FACTOR 0.9994930 GRID FACTOR	389,983.639	S-11
- 4	0 54 00.545	112 23 33.635 CCNVERGENCE ELEVATION	4,457,461.268 0.9997726 SCALE FACTOR 0.9995309 GFID FACTOR	381,570.567	S-12
- (0 54 13.818	112 23 51.558 CCNVERGENCE ELEVATION	4,460,260.113 0.9997737 SCALE FACTOR 0.9995328 GRID FACTOR	381,191.224	S-14
- (0 51 01.851	112 18 52.056 CONVERGENCE ELEVATION	4,463,247.498 0.9997535 SCALE FACTOR 0.9994994 GRID FACTOR	388,309.437	S-15

TOOELE ARMY DEPOT / SOUTH LAMBERT - UTAH CENTRAL

LA	FITUDE	LONGITUDE	NORTHING	EASTING	SAMPLE
40 16 (02.709 (112 21 35.185 CONVERGENCE			I-WZZ
- 0 33 0	09.241 (CONVERGENCE	705,358,800 1,759 0.9999272 SCALE FACTOR 0.9999272 GPID FACTOR	,287:500	SSW-2
- 0 31 2	20,792 C	12 18 56.084 CONVERGENCE LEVATION	703,626.000 1,772 0.9999266 SCALE FACTOR 0.9999266 GPID FACTOR	,395.500	SSD-1
- 0 31 3	9.645 C	12 19 25.515 CONVERGENCE LEVATION	703,551.000 1,770, 0.9999265 SCALE FACTOR 0.9999265 GFID FACTOR	113,200	SSD-2
- 0 33 4	9.923 C	12 22 48.890 ONVERGENCE LEVATION	719,917.600 1,754, 0.9999326 SCALE FACTOR 0.9999326 GRID FACTOR	507,600	SSD-3
- 0 34 3	0.344 C	12 23 51.991 ONVERGENCE LEVATION	723,779.200 1,749, 0.9999342 SCALE FACTOR 0.9999342 GRID FACTOR	657.700	SSD-4

TOOELE ARMY DEPOT / SOUTH U.T.M.- ZONE # 12

_		L	ATITUDE	LONGITUDE	NOFTHING EASTING	SAMPLE #
-	40	16	05.592 44.430	112 21 35.185	4,458,204.034 384,380.681 0.9997645 SCALE FACTOR 0.9997645 GFID FACTOR	SSW-1
-	40	16 52	50.925	112 21 45.382 CONVERGENCE ELEVATION	4,458,040.711 384,137.262 0.9997652 SCALE FACTOR 0.9997652 GFID FACTOR	SSW-2
-	40	15 51	01.100	112 18 56.084 CONVERGENCE ELEVATION	4,457,489.804 388,128.838 0.9997540 SCALE FACTOR 0.9997540 GPID FACTOR	SSD-1
~	4 0 0	15 51	20.196	112 19 25.515 CONVERGENCE ELEVATION	4,457,470.925 387,433.190 0.9997559 SCALE FACTOR 0.9997559 GRID FACTOF	SSD-?
-	40	18 53	J 4 0 Z Z	112 22 48.890 CONVERGENCE ELEVATION	4,462,485.821 382,706.079 0.9997693 SCALE FACTOR 0.9997693 GFID FACTOR	SSD-3
-	4 0 0	19 54	16.156	112 23 51.991 CONVERGENCE ELEVATION	4,463,671.107 381,234.832 0.9997736 SCALE FACTOR 0.9997736 GRID FACTOR	SSD-4

Appendix E

Contamination Results

(Site Summary Forms)



SITE SUMMARY SHEET - TOOELE ARMY DEPOT.

SITE IDENT	TIFICATION	We	11 1-N			
SITE TYPE		Wel	1			
SCREENED INTERVAL 392'-400'; 406'-423'; 434'-470'; 475'-502'; 510'-520'; 573'604';						
630'-640'; 700'-743 DEPTH TO WATER (FEET) 380.6						
GROUND E	LEVATION (FEET) 484	8.1			
LOCATION	793,4	425 Northi	ng, 1,763	.675 East	ing (Utah State Plane, Central Zone)	
					n the North Area; located in	
mair	ntenance a	area, sout	h east qu	arter - s	ampled to obtain background water	
qua!	lity data					
						
						
						
						
						
FIELD SAMPLE NUMBER	SAMPLE DEPTH (FEET)	SAMPLE TYPE	SAMPLE METHOD	DATE SAMPLED	CHEMICAL CONSTITUENTS (ABOVE LOD)	
Well 1	381	ground water	pump	3/30/82	Ni, Zn. Cl. NO ₃ . SO ₄ . Cr. Pb. Na	
		İ				
				<u> </u>		



SITE SUMMARY SHEET - TOOELE ARMY DEPOT.

Zone)

SITE IDENT	TIFICATION .	We]	Well 2-N						
SITE TYPE		We]	Well 390'-480' 389.0						
SCREENED	INTERVAL	390							
DEPTH TO V	NATER (FEE	T)389							
GROUND E	LEVATION (FEET)	55.8 ft.						
				thing 1.	767,875 Easting (Utah State Plane, Centra				
					n North Area; located in maintenance				
area, n	northeast	quarter -	- sampled	to obtain	background water quality data				
area, i	ior dieast	quar cer	sampred	to obtain	background water quality data				
				<u> </u>					
		<u> </u>							
FIELD SAMPLE NUMBER	SAMPLE DEPTH (FEET)	SAMPLE TYPE	SAMPLE METHOD	DATE SAMPLED	CHEMICAL CONSTITUENTS (ABOVE LOD)				
Well 2	391	ground water	pump	3/30/82	Ni, Zn, Cl, NO ₃ , SO ₄ , Cr, Pb, Na				
					50, 50, 50, 504, 50, 10, 50				
		 	 -						
	-		<u> </u>						
<u> </u>									
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SITE SUMMARY SHEET - TOOELE ARMY DEPOT.

SITE IDENTIFICATION			Well 4-N				
			Well				
SCREENED	INTERVAL	638	638'-696'; 700'-775'				
DEPTH TO I	WATER (FEE	T) 622	2				
GROUND E	LEVATION (FEET)	15				
LOCATION		796	300, No	rthing, 1	.731,450 Easting (Utah State Plane Centra	l Zor	
REASON FO	OR SAMPLIN	G <u>Exi</u>	sting Wel	1 No. 4.	North Area; located on ridge near N-6		
in sout	h west qu	arter. S	Sampled to	obtain b	ackground water quality data.		
	····		·····				
							
							
					·		
5.5.5		Ι	T				
FIELD SAMPLE NUMBER	SAMPLE DEPTH (FEET)	SAMPLE TYPE	SAMPLE METHOD	DATE SAMPLED	CHEMICAL CONSTITUENTS (ABOVE LOD)		
Well 4	622	ground water	pump	3/31/82	Ni, Zn, Cl, NO ₃ , SO ₄ , Cr, Pb, Na,		
	-				gross beta		
		-					
	-		<u> </u>				
		-					
			<u> </u>				
1	1	i		i	7		



SITE IDENTIFICATION Well 5-N

SITE TYPE		Wei	11		
SCREENED	INTERVAL				
DEPTH TO V	VATER (FEE	T) 340	<u> </u>		
GROUND EI	LEVATION (FEET)47	50 feet		
LOCATION		809	332.61	Northing,	1,731,456.07 Easting (Utah State Plane
					North Area: located at Firing Range ound water quality data.
FIELD SAMPLE NUMBER	SAMPLE DEPTH (FEET)	SAMPLE TYPE	SAMPLE METHOD	DATE SAMPLED	CHEMICAL CONSTITUENTS (ABOVE LOD)
Well 5	340	ground water	pump	4/14/82	Ni, Zn, Cl, NO ₃ , SO ₄ , Cr, Cu, Pb, Na
Well5-R	340	ground water	pump	6/25/82	
					
	· <u> </u>				



Zone)

SITE IDENT	TIFICATION .	Well	6-N (US	GS-2)	
SITE TYPE		Wel	1		
SCREENED	INTERVAL	375	-415'		
DEPTH TO V	VATER (FEE	T) 247	7.73		
GROUND E	LEVATION (FEET)455	3	<u></u>	
LOCATION		807,584.7	5 Northin	g. 1,752,	526.29 Easting (Utah State Plane Central
REASON FO	OR SAMPLIN	GExi	sting wel	1 No. 6.	USGS well, near east boundary of Ammo
area, r	northeast	quarter;	sampled t	o obtain	background water quality data
					
			···	 	
<u> </u>		_			
<u> </u>					
FIELD SAMPLE NUMBER	SAMPLE DEPTH (FEET)	SAMPLE TYPE	SAMPLE METHOD	DATE SAMPLED	CHEMICAL CONSTITUENTS (ABOVE LOD)
Well 6	248	ground water	pump	4/4/82	Ni, Zn, C1, NO ₃ , SO ₄ , Cr, Pb, Na
 					
					
	<u></u>	<u> </u>	[]		
					
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SITE IDENT	TIFICATION	Wel	Well 1-S								
SITE TYPE Well											
SCREENED	CREENED INTERVAL 290'-404'										
DEPTH TO V	NATER (FEE	T)286	5.71								
GROUND EI	LEVATION (FEET)539	98.4								
LOCATION		727	7, 419 Nor	thing; 1,	770,796 Easting (Utah State Plane, Centra	al Zon					
					South area: located near north boundary ound water quality data.						
FIELD SAMPLE NUMBER	SAMPLE DEPTH (FEET)	SAMPLE TYPE	SAMPLE METHOD	DATE SAMPLED	CHEMICAL CONSTITUENTS (ABOVE LOD)						
Well 1	287	ground water	pump	3/31/82	Ni, Zn, C1, NO3, SO4, Cr, Pb, Na						
	-										
			//								



SITE IDEN	rification.	Wel	1 3-s							
SITE TYPE			·-·	 						
SCREENED	INTERVAL		···							
DEPTH TO	WATER (FEE	T)		· · · · · · · · · · · · · · · · · · ·						
GROUND E	LEVATION (FEET)								
LOCATION										
REASON F	REASON FOR SAMPLINGExisting well No. 3, South area; CAMDS well; sampled to									
<u>obtain</u>	backgroun	<u>id water q</u>	uality da	ta.						
										
				···						
			· · · · · · · · · · · · · · · · · · ·							
5:51.0				<u> </u>						
FIELD SAMPLE NUMBER	SAMPLE DEPTH (FEET)	SAMPLE TYPE	SAMPLE METHOD	DATE SAMPLED	CHEMICAL CONSTITUENTS (ABOVE LOD)					
CAMDS	?	ground water	pump	4/15/82	Ni, Zn, Cl, F, SO ₄ , Cr, Cu, Pb, Na,					
					gross beta					
					-					
		ļ								
										
										
-										



SITE IDENT	TIFICATION	N-:	SDl			
SITE TYPE		dit	ch			
SCREENED	INTERVAL	N/A	<u> </u>			_
DEPTH TO V	NATER (FEE	T) dry	<u> </u>			_
GROUND EI	LEVATION (FEET) <u>not</u>	determir	ned		_
LOCATION		792	2,041.9 No	orthing, 1	,739,734.9 Easting (Utah State Plane,	_ Central
REASON FO	OR SAMPLIN	GSed	liment sam	mple taken	n from dry wash in bottom of flood mainants were migrating by surface wate	_
f]	low from	the chemic	al range	or demoli	tion area.	_
						_
						<u>-</u>
						<u>_</u>
						
		1		γ		_
FIELD SAMPLE NUMBER	SAMPLE DEPTH (FEET)	SAMPLE TYPE	SAMPLE METHOD	DATE SAMPLED	CHEMICAL CONSTITUENTS (ABOVE LOD)	
N-SD1	0.5	sediment	grab	4/3/82	NO ₃ , SO ₄ , Na	
						-
						7
						1
						-



SITE IDENTIFICATION	N-SD2
SITE TYPE	ditch
SCREENED INTERVAL	N/A
DEPTH TO WATER (FEET)	dry
GROUND ELEVATION (FEET)	not taken
LOCATION 813.593.2 N	orthing, 1,743,196.0 Easting (Utah State Plane, Central Zone)
	Sediment sample taken from dry wash along northern boundary
	ontaminants were being carried across boundary by surface
water flow.	
 	

FIELD SAMPLE NUMBER	SAMPLE DEPTH (FEET)	SAMPLE TYPE	SAMPLE METHOD	DATE SAMPLED	CHEMICAL CONSTITUENTS (ABOVE LOD)
N-SD2	0.5	sediment	grab	4/3/82	SO ₄ , Na, As, Zn
	_				



SITE IDENT	TFICATION	N-SI	03							
SITE TYPE		Pone	Pond							
SCREENED INTERVAL N/A										
DEPTH TO WATER (FEET) approx. 6" above sample										
GROUND ELEVATION (FEET) not surveyed										
LOCATION805,676.9 Northing, 1,760,645.9 Easting (Utah State Plane Cent										
REASON FO	R SAMPLIN	GSed:	iment sam	ple taken	from Industrial Waste Pond to	Zone)				
_determi	ne what c	ontaminant	ts precip	itated an	d collected in sediment underlying the	surface				
water.										
				<u> </u>						
	-									
 										
	····			<u> </u>						
	-	1		r		7				
FIELD SAMPLE NUMBER	SAMPLE DEPTH (FEET)	SAMPLE TYPE	SAMPLE METHOD	DATE SAMPLED	CHEMICAL CONSTITUENTS (ABOVE LOD)	<u>}</u>				
N-SD3	0.5	sediment	grab	4/4/82	C1, F, PO ₄ , SO ₄ , Na Ni, Zn	j				
		<u> </u>	i							
	-									
						•				



SITE IDEN	TIFICATION	N-S	D4							
SITE TYPE		Dit	.ch							
SCREENED	CREENED INTERVAL N/A									
DEPTH TO	WATER (FEE	T) dry	,							
GROUND E	LEVATION (FEET) not	determin	ed						
LOCATION		Rad	iological	storage	area					
REASON F	OR SAMPLIN	GSam	pled sedi	ment bein	g carried out of radiological storage					
_and dis	sposal are	a to dete	rmine if	any radio	nuclides were migrating in surface-wate					
_runoff.		_	<u> </u>							
		· 								
										
		Τ								
FIELD SAMPLE NUMBER	SAMPLE DEPTH (FEET)	SAMPLE TYPE	SAMPLE METHOD	DATE SAMPLED	CHEMICAL CONSTITUENTS (ABOVE LOD)					
N-SD4	0.5	sediment	grab	5/26/82	None above LOD					
	 									
]								
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SITE IDEN	TIFICATION	S-:	SD1						
SITE TYPE		di	tch						
SCREENED INTERVAL N/A									
DEPTH TO WATER (FEET)dry									
GROUND E	LEVATION	(FEET) not	t determin	ned					
LOCATION 703,626.0 Northing: 1,772,395.50 Easting (Utah State Plane, Centra									
	OR SAMPLIN				dry wash on southern boundary to	Zone)			
determ	ine if co				surface-water flow from Area 2 or from				
	rning gro								
				·					
									
	γ	T -		Γ					
FIELD SAMPLE NUMBER	SAMPLE DEPTH (FEET)	SAMPLE TYPE	SAMPLE METHOD	DATE SAMPLED	CHEMICAL CONSTITUENTS (ABOVE LOD)				
S-SD1	0.5	sediment	grab	4/4/82	NO3, PO4, SO4, Na As, Zn				
	<u> </u>								
j	-								
		<u> </u>				,			
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Zone)

SITE IDEN	TIFICATION	S-9	SD2		
SITE TYPE		dit	ch		
SCREENED	INTERVAL	N/I	A		
DEPTH TO	NATER (FE	ET)dry	7		
GROUND E	LEVATION ((FEET) Not	determin	<u>red</u>	
LOCATION		703.551.0	Northing	1, 1,770,1	13.20 Easting (Utah State Plane, Central
REASON FO	OR SAMPLING -4 and S	NGSec	liment sam e taken t	mple from co determi	dry wash on southern boundary between ne if contaminants were migrating by
					·
FIELD SAMPLE NUMBER	SAMPLE DEPTH (FEET)	SAMPLE TYPE	SAMPLE METHOD	DATE SAMPLED	CHEMICAL CONSTITUENTS (ABOVE LOD)
S-SD2	0.5	sediment	grab	4/4/82	C1, NO ₃ , SO ₄ , Na, As, Ni, Zn



SITE IDENT	TIFICATION		D3		
SITE TYPE		dit	ch		
SCREENED	INTERVAL	N/A			
LOCATION REASON FO	719.	.917.60 No IGSed	rthing; 1 iment sam	,754,507. mple taken	60 Easting (Utah State Plane, Central Zone) from drainage ditch coming from Area 10 contaminants coming out at Area 10, dead
FIELD SAMPLE NUMBER	SAMPLE DEPTH (FEET)	SAMPLE TYPE	SAMPLE METHOD	DATE SAMPLED	CHEMICAL CONSTITUENTS (ABOVE LOD)
S-SD3	0.5	sediment	grab	4/4/82	C1, NO ₃ , SO ₄ , Na , As, Zn
_					



SITE IDENT	FIFICATION		D4		
SITE TYPE		dit	ch		
SCREENED	INTERVAL	N/A			
DEPTH TO	WATER (FEE	T) dry			
GROUND E	LEVATION (FEET) <u>not</u>	taken	 	
LOCATION	723,779	.20 North	ing; 1,74	9,657.70	Easting (Utah State Plane, Central Zone)
REASON FO	OR SAMPLIN	GSed	iment sam	ple taken	from drainage ditch in northwest quarter.
Sample	taken to	determine	if conta	minants m	igrating across depot by surface-water
		·	-		
FIELD SAMPLE NUMBER	SAMPLE DEPTH (FEET)	SAMPLE TYPE	SAMPLE METHOD	DATE SAMPLED	CHEMICAL CONSTITUENTS (ABOVE LOD)
S-SD4	0.5	sediment	grab	4/4/82	Zn, Cl, SO ₄ , Na _{, As, Ni}
					
			_	İ	



SITE IDENTIFICATION	N-SW1
SITE TYPE	pond
SCREENED INTERVAL	N/A
DEPTH TO WATER (FEET)	surface water
GROUND ELEVATION (FEET)	not taken, ground-surface level
LOCATION 792,581.50 Nort	thing: 1,750,701.40 Easting (Utah State Plane, Central Zone)
REASON FOR SAMPLING	Surface-water sample taken from laundry effluent near TNT
washout pond. This ef	fluent is the mechanism for carrying TNT down through unsat-
urated zone. Sample t	taken to determine contaminants and characterisitics of
this effluent. High v	value in Hazard Ranking System.

FIELD SAMPLE NUMBER	SAMPLE DEPTH (FEET)	SAMPLE TYPE	SAMPLE METHOD	DATE SAMPLED	CHEMICAL CONSTITUENTS (ABOVE LOD)
NSW-1	0.5	surface water	grab	4/5/82	Ni, Zn, RDX, Cl, F, NO ₃ , SO ₄ , Cr, Cu, Ph
NSW1-R	0.5	surface water	grab	6/23/82	Na , lllTCE
					
		<u> </u>		:	



SITE IDENTIFICATION			N-SW2					
			pond					
SCREENED INTERVAL			Α					
DEPTH TO V	VATER (FEE	T) su	rface wate	er				
GROUND EI	EVATION (FEET)not	t taken, a	pprox. 10) below ground surface			
LOCATION	805,676	5.90 North	ning; 1,76	0.645.90	Easting (Utah State Plane, Central Zone)			
	OR SAMPLIN				taken from industrial waste pond directly			
above s	sediment s				ermine contaminants in pond and for			
compar	ison of co	oncentrati	ons found	in water	versus sediment in equilibrium with the			
water.			ard Ranki					
				-				
		<u></u> .	**·					
		,	,	· · · · · · · · · · · · · · · · · · ·				
FIELD SAMPLE NUMBER	SAMPLE DEPTH (FEET)	SAMPLE TYPE	SAMPLE METHOD	DATE SAMPLED	CHEMICAL CONSTITUENTS (ABOVE LOD)			
N-SW2	0.5	surface water	grab	4/14/82	Ni, Zn, TETRYL*, RDX*, CYN, Oil/Grease			
	-				C1, F, NO ₃ , PO ₄ , Be, Cd, Cr, Cu,			
					Pb, Na, 24DMPN			
					* interferences present			



SITE IDENTIFICATION			N-SW3					
SITE TYPE		lag	lagoon					
SCREENED	INTERVAL	N/A	<u> </u>					
DEPTH TO	WATER (FEE	T)sur	face wate	er				
GROUND E	LEVATION (FEET) <u>app</u>	rox. 4665	· · · · · · · · · · · · · · · · · · ·				
LOCATION	_800,252.	.60 Northi	.ng; 1,759	,866.10 E	asting (Utah State Plane, Central Zone)			
					taken from northwest corner of sewage ants in lagoon, because approx. 50 gpm			
		. –			Ranking system.			
				 				
								
	1			·				
FIELD SAMPLE NUMBER	SAMPLE DEPTH (FEET)	SAMPLE TYPE	SAMPLE METHOD	DATE SAMPLED	CHEMICAL CONSTITUENTS (ABOVE LOD)			
N-SW3	0.5	surface water	grab	6/23/82	CYN, C1, PO, SO, Na, Gross beta			
					Zn, Cu phenD6			
			···					
		ŀ						



SITE IDENT	IFICATION	<u>S-S</u>	W1		
SITE TYPE		pon	<u>d</u>		
SCREENED	INTERVAL	N/A			
DEPTH TO V	VATER (FEE	T) sur	face wate	r	
		FEET) <u>not</u>			el
					Easting (Utah State Plane, Central Zon
					taken from crater. Sampled to determin
					urface water runoff. This crater was a
		for a la			
					
					· · · · · · · · · · · · · · · · · · ·
5151.0				<u> </u>	
FIELD SAMPLE NUMBER	SAMPLE DEPTH (FEET)	SAMPLE TYPE	SAMPLE METHOD	DATE SAMPLED	CHEMICAL CONSTITUENTS (ABOVE LOD)
	(1 221)		,,,,,,,,,,	JAN. EED	
S-SW1	0.5	surface water	grab	4/6/82	As Ni 7n Cl NO SO Cr Cu Ph
c cvip	0 "	surface			As, Ni, Zn, C1, NO ₃ , SO ₄ , Cr, Cu, Pb
S-SW1R	0.5	water	grab	6/24/82	Ag, Na, gross alpha, gross beta
					



211E IDEM	TIFICATION	S-	SW2				
SITE TYPE		Su	rface Wat	er			
SCREENED INTERVALN/A							
DEPTH TO	DEPTH TO WATER (FEET) not taken, approx. same as level in Well S-6						
GROUND E	LEVATION (FEET) app	prox. same	as water	c level in S-6		
LOCATION	705,358	.80 North	ing; 1,759	9,287.50 E	Easting (Utah State Plane, Central Zone)		
					taken from standing water in crater near		
Well S	-6. This	water is	probably	hydrologi	ically connected to groundwater since it		
					Well S-6. Sample taken to determine if		
		water and					
			_				
FIELD SAMPLE NUMBER	SAMPLE DEPTH (FEET)	SAMPLE TYPE	SAMPLE METHOD	DATE SAMPLED	CHEMICAL CONSTITUENTS (ABOVE LOD)		
	(FEET)	<u> </u>	WETHOD	SAMPLED	STATISTICAL CONSTITUTION (S (ABOVE LOD)		
S-SW2	0.5	surface water	grab	4/15/82			
		Hacei	grab	4/15/82	Pb, C1, F, SO ₄ , Cd, Cr, Cu, As,		
					Ag, Na, gross alpha		
	 -						
		·					
	ľ						



SITE IDENTIFICATION		S-S	S-SW3						
SITE TYPE			ditch						
SCREENED	INTERVAL	N/A	\						
DEPTH TO	WATER (FEE	T) sur	face wate	r					
GROUND E	LEVATION (FEET) not	taken; a	pproximat	ely ground level				
LOCATION		not	surveyed	l, west si	de of CAMDS				
REASON F	OR SAMPLIN	GSur	face-wate	er effluen	t from CAMDS. Sample taken to determine				
					y close to ground water here. Also,				
numero	us dead ra	abbits dis	covered.	High val	ue in Hazard Ranking System.				
			·						
									
	-				<u> </u>				
	Τ	T		<u> </u>					
FIELD SAMPLE NUMBER	SAMPLE DEPTH (FEET)	SAMPLE TYPE	SAMPLE METHOD	DATE SAMPLED	CHEMICAL CONSTITUENTS (ABOVE LOD)				
s-sw3	0.5	surface water	grab	5/4/82	Zn, C1, NO ₃ , PO ₄ , SO ₄ , Na, Cu				
									
7									
	.		-						
<u> </u>			_						



SITE IDEN	TIFICATION	<u>N-</u>	1	<u> </u>		
SITE TYPE		Su	rf			
SCREENED	INTERVAL	N/	Α			
DEPTH TO	WATER (FE	T)dry	у			
GROUND E	LEVATION ((FEET) no	t taken			
LOCATION						
REASON F	OR SAMPLIN	IGSo:	il sample	taken in	northeast quarter of north area. Sar	mpl
					nich was deleted as unnecessary. This	
sample	measures	contamina	ants in up	per soil	from spill from industrial pond. High	 -
value :	in Hazard	Ranking S	System.			
						_
						_
						_
	,		.			
FIELD SAMPLE NUMBER	SAMPLE DEPTH (FEET)	SAMPLE TYPE	SAMPLE METHOD	DATE SAMPLED	CHEMICAL CONSTITUENTS (ABOVE LOD)	
GS-1	0.5	soil	grab	4/4/82	C1, NO ₃ , PO ₄ , SO ₄ , As, Ni, Zn	
		1				
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		<u>N-</u>							
SCREENED INTERVAL									
				urface					
		T) dry							
		FEET) _46							
LOCATION	806,047	7.12 North	hing; 1,76	0,638.30	Easting (Utah State Plane, Central Zone)				
REASON FO	OR SAMPLIN	GBoi	ring drill	ed for so	il samples and groundwater samples. Soil				
not sui	table for	analysis	. Hit be	drock; no	groundwater. Drilled to sample perched				
		rial pond		· · · · · · · · · · · · · · · · · · ·					
		-							
		·							
				·					
					•				
		<u> </u>		·					
FIELD SAMPLE NUMBER	SAMPLE DEPTH (FEET)	SAMPLE TYPE	SAMPLE METHOD	DATE SAMPLED	CHEMICAL CONSTITUENTS (ABOVE LOD)				
100 0	E TAKEN								
-									
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SITE IDENTIFICATION		N-	N-2B						
SITE TYPE			re						
SCREENED	INTERVAL	ca	pped bore	, no scre	en				
DEPTH TO	WATER (FE	ET)dr	Y						
GROUND E	LEVATION	(FEET) 46	79.19						
LOCATION	806,0	47.12 Nor	thing; 1,	760,638.3	O Easting (Utah State Plane, Central Zone)				
REASON FO	OR SAMPLII	NG Re-	-drilling	of Borin	g N-2A. No samples suitable for collection	'n.			
									
			 						
		<u> </u>							
FIELD SAMPLE NUMBER	SAMPLE DEPTH (FEET)	SAMPLE TYPE	SAMPLE METHOD	DATE SAMPLED	CHEMICAL CONSTITUENTS (ABOVE LOD)				
NO	NE TAKE	V							
									
			· · · · · · · · · · · · · · · · · · ·						



SITE IDENTIFICATION	N-2C
SITE TYPE	well
SCREENED INTERVAL	76.9 to 96.4
DEPTH TO WATER (FEET)	87.9
GROUND ELEVATION (FEET	4681.1
LOCATION <u>805,581.30 N</u>	orthing; 1,760,825 Easting (Utah State Plane, Central Zone)
REASON FOR SAMPLING	Well drilled to sample perched zone coming from seepage of
industrial waste pon	d. Approximately 50 gpm seeping into ground. Very high value
in Hazard Ranking Sy	stem.

FIELD SAMPLE NUMBER	SAMPLE DEPTH (FEET)	SAMPLE TYPE	SAMPLE METHOD	DATE SAMPLED	CHEMICAL CONSTITUENTS (ABOVE LOD)
N-2C	93	ground water	pump	6/24/82	C1, F, PO ₄ , Na, 2,4,6-TNT*
					As, Ni, Zn, Cr, Pb, 12DCLE, T12DCE,
					111TCE, TRCLE
					* interference present
				_	
				i	



SITE IDENTIFICATION	N-3A
SITE TYPE	well
SCREENED INTERVAL	297.8 feet to 337 feet (bottom of well)
	4471.9 feet (from ground surface)
GROUND ELEVATION (FEET)	4726.82 feet (top of protective casing); 4723.9 (ground surface)
LOCATION792,717.36 No.	cthing; 1,750,641.41 Easting (Utah State Plane, Central Zone)
REASON FOR SAMPLING	Well N-3A is located down-gradient of the TNT washout ponds and uent discharge. The TNT washout area was the second most
important area for inv	restigation as determined by the Hazard Ranking System.

FIELD SAMPLE NUMBER	SAMPLE DEPTH (FEET)		SAMPLE METHOD	DATE SAMPLED	CHEMICAL CONSTITUENTS (ABOVE LOD)
GS-1	1.0	sediment	grab	2-27-82	Ni, Zn, Na, 2,4-DNT, 2,6-DNT* NO ₃ , PO ₄ , SO ₄ , 2,4,6-TNT, RDX
ss-3	10.75	soil	split spoon	2-27-82	RDX
SS-5	20.75	soil	split spoon	2-27-82	RDX
SS-7	30.75	soil	split spoon	2-27-82	none above LOD
SS-9	40.75	soil	split spoon	2-27-82	2,4,6-TNT
SS-11	50.75	soil	split spoon	2-27-82	2,4-DNT
SS-13	60.75	soil	split spoon	2-27-82	RDX
SS-15	70.75	soil	split spoon	2-27-82	RDX
SS-17	80.75	soil	split spoon	2-27-82	RDX
SS-19	90.75	soil	split spoon	2-27-82	RDX
SS-22	100.75	soil	split spoon	2-27-82	None above LOD
SS-30	138.75	soil	split spoon	2-27-82	None above LOD
N-3A	2 52 g	roundwater	pump	4-5-82	Ni, Zn, RDX, C1, F, NO ₃ , SO ₄ , Cr, 1

Na, gross beta



SITE IDENT	IFICATION.	N-3	ВВ					_		
SITE TYPE		We]	1							
SCREENED	INTERVAL.	36.	.8 to 56.4	feet						
DEPTH TO V	VATER (FEE	T) 50.	. 9							
GROUND EL	EVATION (I	FEET) <u>472</u>	24.1							
			ng; 1,750	,651.21 E	asting	(Utah	State 1	Plane, C	entral	Zone)
			l drilled						d when	drilling
										
										······································
		_								
		.								
					_					
FIELD	SAMPLE	SAMPLE	CANADI E							

FIELD SAMPLE NUMBER	SAMPLE DEPTH (FEET)	SAMPLE Type	SAMPLE METHOD	DATE SAMPLED	CHEMICAL CONSTITUENTS (ABOVE LOD)
N-3B	59:9 to	ground water	bailed	6-23-82	C1, F, NO ₃ , SO ₄ , Na, Gross beta
					As, Zn, Cr, Cu, 46DN2C



SITE IDEN	TIFICATION	N-4	<u> </u>	·					
SITE TYPE	TYPE Well								
SCREENED	CREENED INTERVAL 193.1 to 208 feet								
DEPTH TO	WATER (FEE	T) 191	.3 feet	··· <u>-</u>					
GROUND E	LEVATION (FEET)466	52.6						
LOCATION	_801.178	63 Northi	ng; 1,759	,476.34 E	asting (Utah State Plane, Central Zone)				
					200 feet north-northwest of the sewage				
lagoon	. It is r	re-located	and comb	ined with	N-1. Sampled to determine if contami-				
nants	in ground-	-water fro	m sewage	lagoon an	d landfill sources. High value in				
Hazard	Ranking S	System.							
FIELD SAMPLE NUMBER	SAMPLE DEPTH (FEET)	SAMPLE TYPE	SAMPLE METHOD	DATE SAMPLED	CHEMICAL CONSTITUENTS (ABOVE LOD)				
N-4	196	ground		1					
		water	pump	6/24/82	C1, F, NO ₂ , SO ₄ , Na, Gross beta				
		water	pump	6/24/82	C1, F, NO ₃ , SO ₄ , Na, Gross beta				
		water	pump	6/24/82	C1, F, NO ₃ , SO ₄ , Na, Gross beta Ni, Zn, TRCLE				
		water	pump	6/24/82	<u> </u>				
		water	pump	6/24/82	<u> </u>				
		water	pump	6/24/82	<u> </u>				
		water	pump	6/24/82	<u> </u>				
		water	pump	6/24/82	<u> </u>				
		water	pump	6/24/82	<u> </u>				
		water	pump	6/24/82	<u> </u>				
		water	pump	6/24/82	<u> </u>				
		water	pump	6/24/82	<u> </u>				



SITE IDENT	IFICATION.	N-0	5		
SITE TYPE		We:	L1		
SCREENED	INTERVAL	63:	3.1 to 692	feet	
DEPTH TO V	VATER (FEE	T) dry	Z		
GROUND EL	_EVATION (F	EET)509	91.9		
				,581.60 E	asting (Utah State Plane, Central Zone)
					ge overlooking chemical range. Drilled
					d demolition range. Well dry- drilled
to 704					
-					
FIELD SAMPLE NUMBER	SAMPLE DEPTH (FEET)	SAMPLE TYPE	SAMPLE METHOD	DATE SAMPLED	CHEMICAL CONSTITUENTS (ABOVE LOD)
N	ONE TAKEN				
					
					
	1				



SITE IDEN	TIFICATION .	N-	7		
SITE TYPE		bor	ce		
SCREENED	INTERVAL.	car	oped bore,		
DEPTH TO	WATER (FEE	T) dry	<u> </u>		
LOCATION	787,720	.53 North	ning; 1,75	7,957.54	Easting (Utah State Plane, Central Zone
REASON FO	OR SAMPLIN	G <u>Wel</u>	ll N-7 was	drilled	on the southern boundary to determine
					ndary from chemical operations near
Stockto	n Bar. W	ell not c	complete -	low prio	rity.
			<u> </u>		
		-			
		r 		¥	
FIELD SAMPLE NUMBER	SAMPLE DEPTH (FEET)	SAMPLE TYPE	SAMPLE METHOD	DATE SAMPLED	CHEMICAL CONSTITUENTS (ABOVE LOD)
	NONE TAKE	И			
			<u> </u>		
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SITE TYPE well SCREENED INTERVAL 53.9 to 73.6 feet DEPTH TO WATER (FEET) dry GROUND ELEVATION (FEET) 4471.9 LOCATION 813.744.46 Northing; 1,752,850.25 Easting (Utah State Plane, Central Zone) REASON FOR SAMPLING Well N-8A drilled along north boundary to locate possible percapher possibly resulting from Industrial Waste Pond. Well dry - no perched zone. FIELD SAMPLE SAMPLE SAMPLE METHOD SAMPLED CHEMICAL CONSTITUENTS IABOVE LOD! NONE TAKEN						
SCREENED INTERVAL 53.9 to 73.6 feet DEPTH TO WATER (FEET) dry GROUND ELEVATION (FEET) 4471.9 LOCATION 813,744.46 Northing; 1,752,850.25 Easting (Utah State Plane, Central Zone) REASON FOR SAMPLING Well N-8A drilled along north boundary to locate possible percapone possibly resulting from Industrial Waste Pond. Well dry - no perched zone. FIELD SAMPLE SAMPLE SAMPLE SAMPLE DATE NUMBER (FEET) SAMPLE METHOD DATE SAMPLED CHEMICAL CONSTITUENTS (ABOVE LOD)						
GROUND ELEVATION (FEET)	SITE TYPE		we	11		
GROUND ELEVATION (FEET) 4471.9 LOCATION 813.744.46 Northing; 1,752,850.25 Easting (Utah State Plane, Central Zone) REASON FOR SAMPLING Well N-8A drilled along north boundary to locate possible percentage and possibly resulting from Industrial Waste Pond. Well dry - no perched zone. FIELD SAMPLE DEPTH SAMPLE SAMPLE SAMPLE DATE CHEMICAL CONSTITUENTS (ABOVE LOD)	SCREENE	INTERVAL	53	.9 to 73.	6 feet	
LOCATION 813,744.46 Northing; 1,752,850.25 Easting (Utah State Plane, Central Zone) REASON FOR SAMPLING Well N-8A drilled along north boundary to locate possible percentage and possibly resulting from Industrial Waste Pond. Well dry - no perched zone. FIELD SAMPLE DEPTH SAMPLE SAMPLE DEPTH NUMBER (FEET) SAMPLE TYPE METHOD SAMPLED CHEMICAL CONSTITUENTS (ABOVE LOD)	DEPTH TO	WATER (FE	ET) dr	У		
REASON FOR SAMPLING Well N-8A drilled along north boundary to locate possible percape zone possibly resulting from Industrial Waste Pond. Well dry - no perched zone. FIELD SAMPLE DEPTH SAMPLE SAMPLE DATE SAMPLE DATE CHEMICAL CONSTITUENTS (ABOVE LOD)	GROUND E	LEVATION	(FEET) <u>44</u>	71.9		
REASON FOR SAMPLING Well N-8A drilled along north boundary to locate possible percape zone possibly resulting from Industrial Waste Pond. Well dry - no perched zone. FIELD SAMPLE DEPTH SAMPLE SAMPLE DATE SAMPLE DATE CHEMICAL CONSTITUENTS (ABOVE LOD)	LOCATION	813,744	.46 North	ing; 1,75	2,850.25	Easting (Utah State Plane Control C
FIELD SAMPLE DEPTH SAMPLE DATE METHOD SAMPLED CHEMICAL CONSTITUENTS (ABOVE LOD)	REASON F	OR SAMPLIN	NG We	11 N-8A d	rilled al	ong north hour days de la lettral zone)
FIELD SAMPLE SAMPLE DATE SAMPLE TYPE METHOD SAMPLED CHEMICAL CONSTITUENTS (ABOVE LOD)	zone p	ossibly r	esulting	from Indu	strial Wa	ste Pond Well dry - no nearly !
SAMPLE DEPTH SAMPLE SAMPLE DATE SAMPLED CHEMICAL CONSTITUENTS (ABOVE LOD)						no perched zone.
SAMPLE DEPTH SAMPLE SAMPLE DATE SAMPLED CHEMICAL CONSTITUENTS (ABOVE LOD)						
SAMPLE DEPTH SAMPLE TYPE METHOD SAMPLED CHEMICAL CONSTITUENTS (ABOVE LOD)						
SAMPLE DEPTH SAMPLE SAMPLE METHOD SAMPLED CHEMICAL CONSTITUENTS (ABOVE LOD)						
SAMPLE DEPTH SAMPLE SAMPLE DATE SAMPLED CHEMICAL CONSTITUENTS (ABOVE LOD)						
SAMPLE DEPTH SAMPLE SAMPLE METHOD SAMPLED CHEMICAL CONSTITUENTS (ABOVE LOD)						
SAMPLE DEPTH SAMPLE SAMPLE DATE SAMPLED CHEMICAL CONSTITUENTS (ABOVE LOD)						
SAMPLE DEPTH SAMPLE SAMPLE DATE SAMPLED CHEMICAL CONSTITUENTS (ABOVE LOD)		T -	1			
NOMBER (FEET) TYPE METHOD SAMPLED CHEMICAL CONSTITUENTS (ABOVE LOD)	SAMPLE	DEPTH		SAMPLE	DATE	
NONE TAKEN NONE TAKEN NONE TAKEN	NUMBER	(FEET)	TYPE			CHEMICAL CONSTITUENTS (ABOVE LOD)
NONE TAKEN			j		 	
		NONE TA	KEN		 	
					 	
			·			
		}	1	T		



SITE IDEN	TIFICATION	N-	8B		
SITE TYPE		we	11		
SCREENED	INTERVAL	24	0.73 to 2	80.0 feet	, 181.76 to 201.37 feet
		ET) 16			
GROUND E	LEVATION ((FEET)44	72.3		
LOCATION	<u>813,74</u>	4.61 Nort	hing; 1,7	52,833.47	Easting (Utah State Plane, Central Zone
REASON F	OR SAMPLIN	IGWe	<u>ll N-8B i:</u>	s on nort	hern boundary. This well was drilled
to sam	ple water	leaving l	boundary a	at Depot :	in area most likely to be contaminated.
Two se	parate sc	reens plac	ced to sar	mple two	possibly separate aquifers.
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			· · · · · · · · · · · · · · · · · · ·		
FIELD SAMPLE NUMBER	SAMPLE DEPTH (FEET)	SAMPLE TYPE	SAMPLE METHOD	DATE SAMPLED	CHEMICAL CONSTITUENTS (ABOVE LOD)
N-8B-1	190	ground water	pump	5/3/82	Ni, Zn, Oil/Grease, Cl, NO ₃ , SO ₄ , Na _{,C}
N-8B-2	260	ground water	pump	5/3/82	Ni, Zn, C1, NO ₃ ; SO ₄ , Na, Cr
					
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T					



SITE IDENTIFICATION	S-1
SITE TYPE	well
SCREENED INTERVAL	10.3 to 20.2 feet
DEPTH TO WATER (FEET)	8.0 feet
GROUND ELEVATION (FEET)	approx. 5038
LOCATION711,679.69 No.	rthing; 1,758,523.04 Easting (Utah State Plane, Central Zone)
REASON FOR SAMPLING	Well S-1 is located downgradient from the CAMDS facility led to determine if any contaminants are entering the
groundwater from the	effluent and if any previous contamination is present and being
	nd-water. High value in Hazard Ranking System.

FIELD SAMPLE NUMBER	SAMPLE DEPTH (FEET)	SAMPLE TYPE	SAMPLE METHOD	DATE SAMPLED	CHEMICAL CONSTITUENTS (ABOVE LOD)
SS-2	5.8	soil	splitspoor	5/27/82	Cl, SO4, Na, As, Ni, Zn, DEP
SS-3	10.6		splitspoor		
SS-4	15.6	soil	splitspoor	5/27/82	
SS-5b	20.7	soil	splitspoon	5/27/82	C1, F, SO ₄ , Na, As, Zn, DEP
S-1	12	ground water	pump	6/24/82	Be, Cu. Na. Cr
S-1dup	12	ground water	amuq	6/24/82	



SITE IDENTIFICATION	S-2
SITE TYPE	well
SCREENED INTERVAL	·
DEPTH TO WATER (FEET)	57.8 feet
GROUND ELEVATION (FEET)	5145.6
LOCATION	thing; 1,770,461.39 Easting (Utah State Plane, Central Zone)
REASON FOR SAMPLING	Well S-2 ranked high in our Hazard Ranking System because
of a no-longer-existing	g pond and Bldg. T-600 where there was a very high probability
of contamination.	

FIELD SAMPLE NUMBER	SAMPLE DEPTH (FEET)	SAMPLE TYPE	SAMPLE METHOD	DATE SAMPLED	CHEMICAL CONSTITUENTS (ABOVE LOD)
SS-2	5.5	soil	splitspoo	n 2/19/82	Zn, C1, NO ₄ ,PHEND6 Na
SS-6A	25.3	soil	splitspoo	n 2/19/82	
SS-10	45.8	soil	splitspoo	n 2/19/82	
S-2	65	ground water	pump	4/29/82	Zn, Cl, NO ₃ , SO ₄ , Na, gross alpha, Cr
					3 4 , CI



SITE IDENTIFICATION	S-3
SITE TYPE	well
SCREENED INTERVAL	24.0 - 43.6 feet
DEPTH TO WATER (FEET)	26.85 feet
GROUND ELEVATION (FEET)	5051.3
	thing; 1,756,677.58 Easting (Utah State Plane, Central Zone)
Groundwater was sample pit inside Area 10 whe	Well S-3 is located on the south boundary of Area 10. d to determine if contaminants were leaching from the gravel re mustard was burned and buried. High value in

FIELD SAMPLE NUMBER	SAMPLE DEPTH (FEET)	SAMPLE TYPE	SAMPLE METHOD	DATE SAMPLED	CHEMICAL CONSTITUENTS (ABOVE LOD)
S-3	32	ground water	pump	4/29/82	As, Ni, Zn, C1, F, SO ₄ , Na, gross beta



SITE IDENT	IFICATION.	s-	-4		
SITE TYPE		We	11		
SCREENED	INTERVAL.	64	.0 to 83.	6 feet	
DEPTH TO V	VATER (FEE	T) 58	3.3		
•	EVATION (064.0		
LOCATION	703,646.	17 Northi	ng; 1,770	0,955.03 Eas	ting (Utah State Plane, Central Zone)
REASON FO	R SAMPLIN	GWe	11 S-4 is	s located on	the southern boundary due south of
the_bu	rning or	demilitar	ization a	area. Sampl	es taken from Well S-4 monitor the
ground	water imm	mediately	before le	eaving the d	epot. High value in Hazard Ranking
System	1				
					
	· · · · · · · · · · · · · · · · · · ·				
FIELD SAMPLE NUMBER	SAMPLE DEPTH (FEET)	SAMPLE TYPE	SAMPLE METHOD	DATE SAMPLED	CHEMICAL CONSTITUENTS (ABOVE LOD)

FIELD SAMPLE NUMBER	SAMPLE DEPTH (FEET)	SAMPLE TYPE	SAMPLE METHOD	DATE SAMPLED	CHEMICAL CONSTITUENTS (ABOVE LOD)
S-4	58.4 to 61.0	ground water	bail	5/2/82	A _s , Zn, C1, F, SO ₄ , Na, gross beta



SITE IDENT	TIFICATION.	S-	-5		
SITE TYPE			11		
SCREENED	INTERVAL.	35	7_6_to_57.	6 feet	
DEPTH TO V	NATER (FEE	T) 71	.5 feet		
GROUND EI	LEVATION (I	FEET)50	48.6		
LOCATION	703,623.	07 Northi	ng; 1,768	,779.69 E	asting, (Utah State Plane, Central Zone)
REASON FO	OR SAMPLIN	GWe	11 S-5 is	located	on the southern boundary due south of th
<u>burnir</u>	ng or demi	ilitarizat	ion area.	Samples	taken from Well S-5 monitor the ground
water	immediate	ely before	leaving	the depot	. High value in Hazard Ranking System.
	<u> </u>				
				·	
					
FIELD	SAMBLE			<u> </u>	
SAMPLE NUMBER	SAMPLE DEPTH (FEET)	SAMPLE TYPE	SAMPLE METHOD	DATE SAMPLED	CHEMICAL CONSTITUENTS (ABOVE LOD)
s-5	42	ground water	pump	5/2/82	No. 7- 01 D NO. 00 H
		Water	ратр	3/2/82	As, Zn, C1, F, NO ₃ , SO ₄ , Na, gross bet
·	·		· · · · · · · · · · · · · · · · · · ·		
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SITE IDENTIFICATION _____S-6

SITE TYPE		we	11			
SCREENED	INTERVAL	15	.25 to 34	.93 feet		
DEPTH TO V	VATER (FEE	T) 17	7.3 feet			
GROUND EI	LEVATION (F EET)50	36.9			
LOCATION	705,205	.60 Northi	ing; 1,759	,265.33 E	Casting (Utah State Plane, Central Zone)	
					among the high explosive test craters	
	· · · · · · · · · · · · · · · · · · ·	Ranking		entering	the ground water from these sources.	High
value	In Hazard	Ranking	System.			
						
				 -		
FIELD SAMPLE NUMBER	SAMPLE DEPTH (FEET)	SAMPLE TYPE	SAMPLE METHOD	DATE SAMPLED	CHEMICAL CONSTITUENTS (ABOVE LOD)	
S-6	25	ground water	pump	4/28/82	As, Zn, C1, F, SO ₄ , Na, gross beta	
	:					
			<u></u>			



SITE IDENTIFICATION	s-7
	well
	32.0 to 51.61 feet
	26.5 feet
•	5045_9
	rthing: 1,762,771.55 Easting (Utah State Plane Central Zone)
	Well S-7 is located among the high explosive-test craters
and also downdradient	from the windrows where explosives and other materials were
burned and buried. H	igh value in Hazard Ranking System.

FIELD SAMPLE NUMBER	SAMPLE DEPTH (FEET)	SAMPLE TYPE	SAMPLE METHOD	DATE SAMPLED	CHEMICAL CONSTITUENTS (ABOVE LOD)
S-7	35	ground water	pump	5/4/82	As, Ni, Zn, Cl, F, NO ₃ , SO ₄ , Na, gross beta



SITE IDENTIFICATION	S-8
SITE TYPE	well
SCREENED INTERVAL	65.0 to 84.6 feet
DEPTH TO WATER (FEET)	73.9 feet
GROUND ELEVATION (FEET)	5187.4
LOCATION716,538.06 Nor	thing; 1,771,207.81 Easting (Utah State Plane, Central Zone)
REASON FOR SAMPLING	Well S-8 is located south of Bldg. 553. It was drilled
to determine if contar	mination has occurred from the operations from Bldgs. 554
and 553, the holding	oond (dry) at Bldg 554, and the current effluent from Bldg.
553.	

FIELD SAMPLE NUMBER	SAMPLE DEPTH (FEET)	SAMPLE TYPE	SAMPLE METHOD	DATE SAMPLED	CHEMICAL CONSTITUENTS (ABOVE LOD)
SS-4_	15.7	soil	splitspoo	n 3/2/82	C1, SO ₄ , PHEND6, Na, As, Ni, Zn
SS-9	40.8	soil	splitspoor	3/2/82	Cl, F, NO ₃ , SO ₄ , Na, As, Ni, Zn
SS-12	55.8	soil	splitspoo		
SS-14	65.5	soil	splitspoo	1 3/2/82	
SS-17	80.5	soil	splitspoo	3/2/82	
S-8	80	ground water	ритр	5/4/82	7 311
				<u> </u>	



SITE IDENTIFICATION			S- 9								
SITE TYPE			bore								
SCREENED INTERVAL			capped bore - no screen								
DEPTH TO	WATER (FEE	T) <u>dı</u>	dry								
GROUND E	LEVATION (FEET)50									
LOCATION	706,750	0.40 North	thing: 1,770,801.65 Easting (Utah State Plane, Central Zone)								
REASON F	OR SAMPLIN	GBo	oring S-9	was origi	inally drilled to monitor ground water						
					a 2, to determine the background level						
of co	ntaminants	entering	this are	a. Reach	ned 110 feet and discontinued drilling be	ecause					
did n	ot reach o	ground wat	er.								
		-									
											
		·									
	T		 .								
FIELD SAMPLE NUMBER	SAMPLE DEPTH (FEET)	SAMPLE TYPE	SAMPLE METHOD	DATE SAMPLED	CHEMICAL CONSTITUENTS (ABOVE LOD)						
	NONE TAK	EN									
·											
						•					



SITE IDENTIFICATION	S-10
SITE TYPE	well
SCREENED INTERVAL	70.0 to 89.6 feet
DEPTH TO WATER (FEET)	67.7 feet
GROUND ELEVATION (FEET)	5122.5
LOCATION _ 710,554.72 Nort	thing; 1,772,880.39 Easting (Utah State Plane, Central Zone)
REASON FOR SAMPLING	Well S-10 was drilled downgradient of Area 2 and several
chemical and agent sp	ills to determine if any contamination occurred from these
sources. High rank in	n Hazard Ranking System.

FIELD SAMPLE NUMBER	SAMPLE DEPTH (FEET)	SAMPLE TYPE	SAMPLE METHOD	DATE SAMPLED	CHEMICAL CONSTITUENTS (ABOVE LOD)
S-10	75	ground water	рштр	5-2-82	Zn, C1, NO ₃ , SO ₄ , Na, Cr



SITE IDENTIFICATION			S-11				
SITE TYPE			ore				
SCREENED	INTERVAL	g:	routed to				
DEPTH TO	WATER (FEE	T)d:	ry				
		FEET)5					
					asting (Utah State Plane, Central Zone)		
					led to monitor the unsaturated zone near		
					contaminants were migrating towards		
the wa	ater table	e					
	 						
					·		
	,						
FIELD SAMPLE NUMBER	SAMPLE DEPTH (FEET)	SAMPLE TYPE	SAMPLE METHOD	DATE SAMPLED	CHEMICAL CONSTITUENTS (ABOVE LOD)		
CS-1	17.5	soil	splitspoor	1 2/22/82	C1, NO ₃ , SO ₄ , PHEND6, Zn. Na		
CS_2	37.0			_,,	Zn, Na		

	FIELD SAMPLE NUMBER	SAMPLE DEPTH (FEET)	SAMPLE TYPE	SAMPLE METHOD	DATE SAMPLED	CHEMICAL CONSTITUENTS (ABOVE LOD)
-	CS-1	17.5	soil	splitspoor	1 2/22/82	C1, NO ₃ , SO ₄ , PHEND6, Zn. Na
-	CS-2	37.0	soil	splitspoor	1 2/22/82	
\mid	SS-16	76.2	soil	splitspoo	2/22/82	
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-						
-						
-						
+						
F						
-						
-						
L					<u> </u>	



SITE IDENTIFICATION			S-12					
SITE TYPE			well					
SCREENED	INTERVAL	3	34.0 to 38.9 feet					
DEPTH TO	WATER (FEE	ET)3	1.0 feet					
GROUND E	LEVATION (FEET)5	051.1					
LOCATION	_703,409	15 North	ing: 1.750	0.876.23	Easting (Utah State Plane Central Zone)			
REASON F	OR SAMPLIN	IGW	ell S-12,	original:	ly designed as a cluster well, monitors			
groun	dwater fl	ow off the	e depot at	t the most	likely site of contamination from			
sever	al upgrad	ient sour	ces includ	ding crate	ers and CAMDS. Only one well drilled du			
to co	st constr	aints. Al	lso, Well	S-12 is m	possibly in a ground water discharge are			
				<u> </u>	a ground water discharge are			
	· · · · · · · · · · · · · · · · · · ·							
		 -						
FIELD	SAMPLE	T	<u> </u>	T				
SAMPLE NUMBER	DEPTH (FEET)	SAMPLE TYPE	SAMPLE METHOD	DATE SAMPLED	CHEMICAL CONSTITUENTS (ABOVE LOD)			
		<u> </u>	<u> </u>					
S-12	30	ground	bailed	4/28/82	As, C1, F, NO3, SO4, Na, gross beta			
					na, gross beta			
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SITE IDENTIFICATION S-14
SITE TYPEwell
SCREENED INTERVAL 12.0 to 31.6 feet
DEPTH TO WATER (FEET) 10.65 feet
GROUND ELEVATION (FEET) 5035.9
LOCATION 712,585.81 Northing; 1,749,578.90 Easting (Utah State Plane, Central Zone
REASON FOR SAMPLING Well S-14 was drilled on the west boundary to monitor
groundwater flow going off the depot downgradient of sources such as Area 10 and
possibly CAMDS. Well S-14 is possibly in a groundwater discharge area.
FIELD SAMPLE SAMPLE DATE

FIELD SAMPLE NUMBER	SAMPLE DEPTH (FEET)	SAMPLE TYPE	SAMPLE METHOD	DATE SAMPLED	CHEMICAL CONSTITUENTS (ABOVE LOD)
S-14	15	ground water	pump	5/2/82	As, Ni, Zn, Oil/Grease, Cl, F, SO ₄ , Na, gross beta



SITE IDENTIFICATION			S-15						
SITE TYPE		bo	bore grouted to surface						
SCREENED	INTERVAL	gı							
			dry						
GROUND ELEVATION (FEET) 5311.8									
LOCATION 722,522.46 Northing; 1,772,879.80 Easting (Utah State Plane Central Zone									
REASON F	OR SAMPLIN	IGBo	oring S-15	was dril	led to determine if contaminants were				
leach	ing from t	the old la	ndfill so	urce. No	samples suitable for analysis were				
colle									
			-						
				·					
									
				· · · · · · · · · · · · · · · · · · ·					
FIELD SAMPLE NUMBER	SAMPLE DEPTH (FEET)	SAMPLE TYPE	SAMPLE METHOD	DATE SAMPLED	CHEMICAL CONSTITUENTS (ABOVE LOD)				
	NONE TAK	EN							
				<u> </u>					
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···									